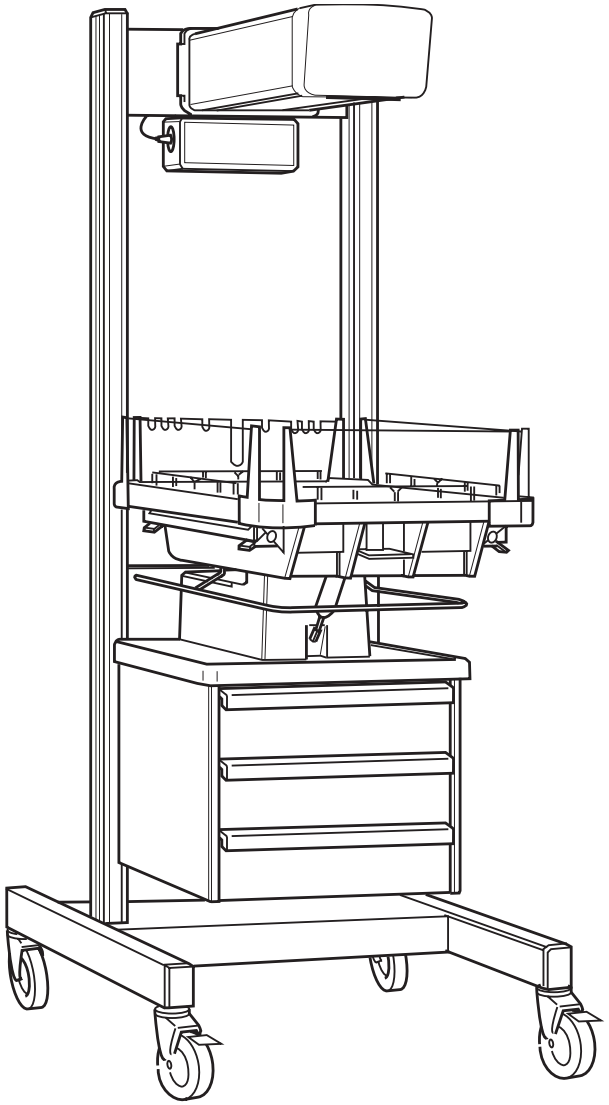
OHMEDA Medical

Ohio® Infant Warmer Systems

All Models

Operation and Maintenance Manual



User Responsibility

This Product will perform in conformity with the description thereof contained in this operating manual and accompanying labels and/or inserts, when assembled, operated, maintained and repaired in accordance with the instructions provided. This Product must be checked periodically. A defective Product should not be used. Parts that are broken, missing, plainly worn, distorted or contaminated should be replaced immediately. Should such repair or replacement become necessary, Ohmeda Medical recommends that a telephone or written request for service advice be made to the nearest Ohmeda Medical Regional Service Center. This Product or any of its parts should not be repaired other than in accordance with written instructions provided by Ohmeda Medical and by Ohmeda Medical trained personnel. The Product must not be altered without the prior written approval of Ohmeda Medical's Quality Assurance Department. The user of this Product shall have the sole responsibility for any malfunction which results from improper use, faulty maintenance, improper repair, damage, or alteration by anyone other than Ohmeda Medical.

CAUTION: U.S. Federal law restricts this device to sale by or on the order of a licensed medical practitioner.

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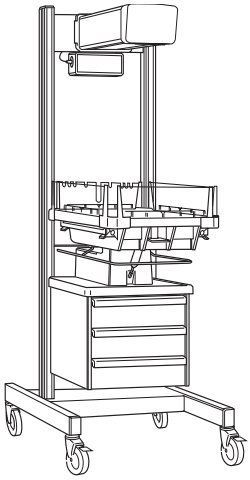
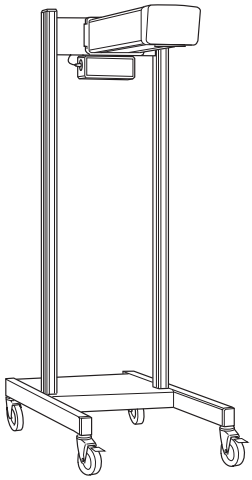
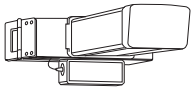


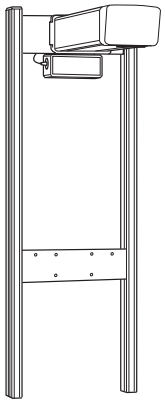
3000 IWS



3050 IWS

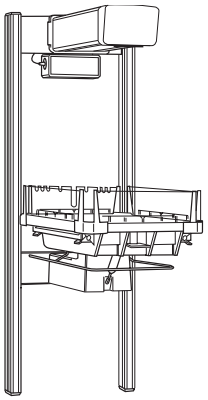


3100 IWS

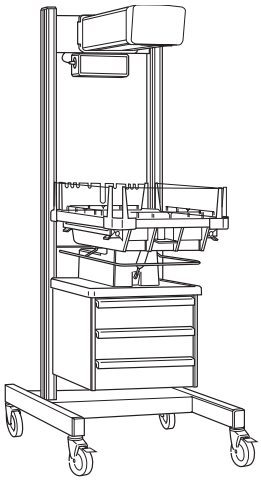


CI.11.003, CI.05.071, CI.05.072, CI.05.073

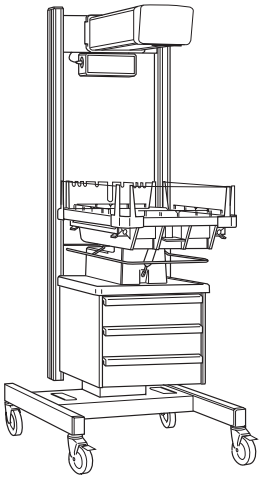
3150 IWS



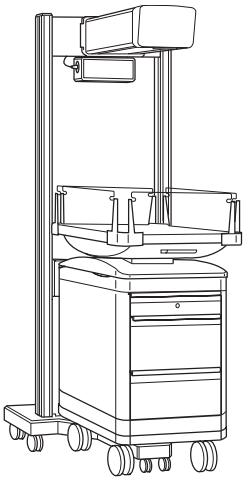
3300 IWS



3400 IWS

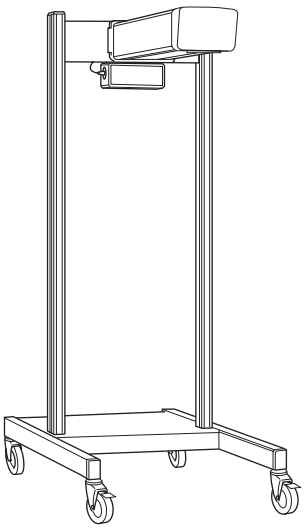


3500 IWS

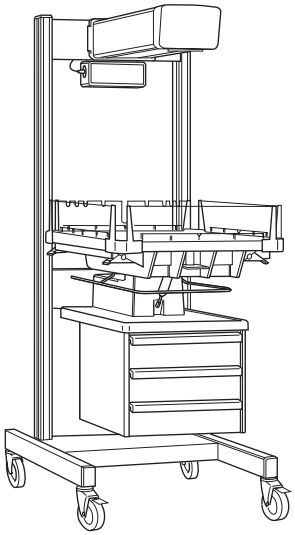


CI.05.074, CI.11.003, CI.05.075, CI.10.022

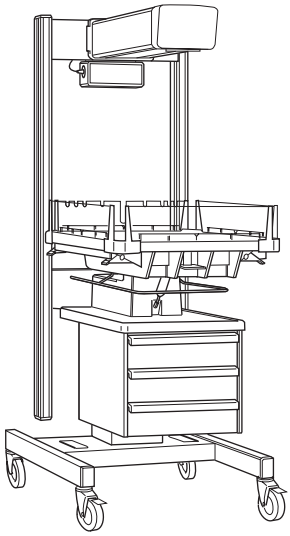
4000 IWS



4300 IWS



4400 IWS



CI.13.004, CI.13.005, CI.13.006

Model Descriptions

All models of the Ohmeda Medical Infant Warmer System provide a controlled source of radiant heat for infants and pediatric patients. The control system uses a microprocessor and provides both manual and servo modes of operation (except for the 2001 International, which is manual mode only).

The model 2001 International has an integral infant bed and is a manual mode only warmer designed for short term attended care in the OR and L&D.

The wall mounted 3050, 3100 and 3150 models are available to accommodate specific architectural requirements in the NICU, L&D, LDR and LDRP room . The 3050 is a heater head only, the 3100 is a heater and dove tail rails, and the 3150 includes a heater, rails and integral bed.

The model 3300 has an integral bed for infants and is intended for procedures and long term care in the Nursery and L&D The model 3400 is the same as the 3300, except it has an elevating pedestal to raise and lower the bed .

The model 3000, 3500 and 4000 are free standing Warmers which can be used over a variety of infant bassinets in the general nursery, over post-operative patients, during patient feeding, and in any other application where controlled radiant heat is required.

The 3500 System also has a detachable Bassinet for infants, and is intended for procedures and long term care in the Nursery, L&D, LDR and LDRP Rooms.

The model 4300 and 4400 have a larger size bed than the model 3300 and are intended for surgical procedures and long term care in the Nursery and L&D Rooms. The model 4400 has an elevating pedestal to raise and lower the bed-to-floor height. The model 4300 is a non-elevating version of the 4400.

IWS Model Options

	Standard Heater/Bed			**Large Heater/Bed**
	2000 Series	**3000 Series**	**3500 Series**	**4000 Series**
Free Standing		3000	3500†	4000
Wall mounted, heater head only		3050		
Wall mounted heater with rails		3100		
Wall mounted heater with rails and bed		3150		
Free standing with integral bed	2001	3300		4300
Elevating pedestal		3400		4400

† Detachable bassinet and wood grain finish standard.

Definitions

Note: A Note provides additional information to clarify a point in the text.

Important: An Important statement is similar to a note but used for greater emphasis.

CAUTION: A Caution statement is used when the possibility of damage to the equipment exists.

WARNING: A Warning statement is used when the possibility of injury to the patient or the operator exists.

Symbols

Attention! Consult the manual for more information.

Type B equipment.

Functional Earth Terminal

Protection Earth Terminal

Increase % Power or Control Temperature

Decrease % Power or Control Temperature

Alarm Silence

Alternating Current

Do not place an infant on this surface

Notes

General Precautions

⚠Warnings

Before using the Ohmeda Medical Infant Warmer System, read this entire manual. Attempting to use this device without a thorough understanding of its operation may result in patient or user injury. This device should only be operated by personnel trained in its operation and under the direction of qualified medical personnel familiar with the risks and benefits of this type of device. **Additional precautions specific to certain procedures are found in the text of this manual.**

Do not use the Warmer in the presence of flammable anesthetics; a possible explosion hazard exists under these conditions.

Do not perform the Checkout Procedures (Mechanical and Control Unit) while a patient occupies the Warmer.

Complete the “Checkout Procedures” section of this manual before putting the unit into operation. If the warmer fails any portion of the checkout procedures it must be removed from use and repaired.

Do not use the Warmer system if the system failure alarm is activated. Remove the unit from service and refer to qualified personnel for repair.

⚠Cautions

Only competent individuals trained in the repair of this equipment should attempt to service it as detailed in the service manual. The Service Manual provides detailed information solely for use by individuals having proper knowledge, tools and test equipment, and for service representatives trained by Ohmeda Medical.

Notes

1/Description

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1.1 General

All models of the Ohmeda Medical Infant Warmer System provide a controlled source of radiant heat for infants and pediatric patients. The control system uses a microprocessor and provides both manual and servo modes of operation (model 2001 International is manual mode only). The patient temperature, control temperature, and elapsed time displays are digital for ease of viewing. The control panel includes a one hour elapsed timer with time displayed in minutes and seconds. The timer features optional audible tones for use during Apgar scoring. A complete audio and visual alarm system is included on the control panel.

The optional ThermaLink feature simplifies charting and monitoring. RS-232 serial data is provided for use with customized research software or compatible vital signs monitors. The Nurse Call connection lets you integrate the Warmer into your current or future remote alarm systems.

1.2 Support Structure

The support structure provides excellent stability for the radiant heater and optional accessories. On free standing units, the support structure consists of the base assembly and the uprights with an integral rail system. The rails provide a means for mounting accessories and ancillary equipment to the unit. Refer to Section 1.7 for a description of the accessory items.

1.3 Heater Assembly

The warmer is designed to warm infants on a bed surface located beneath the heater assembly.

WARNING: ⚠ **Bed-to-heater spacing less than the recommended 64 cm will result in incorrect operation and may adversly affect the patient's condition.**

For your convenience, on Model 3000, 3100, 3500, and 4000 the proper bed-to-heater spacing is indicated by a label located on the right aluminum upright (as viewed from the front).

Warmers shipped since 1993 have high wattage heaters that allow bed to heater spacing of 69 + 5 - 23 cm, making them compatible with many kinds of bassinets.

Since very low birth weight babies may require tighter control of the heat provided to them, we recommend that on warmer models that incorporate a built-in bed and are intended for care of low birth weight babies, the distance between heater and bed be maintained at 69±5 cm.

The heater assembly consists of a radiant heater, parabolic reflector, observation light, and a visual alarm light. The parabolic reflector focuses radiant energy on the bed surface, minimizing energy loss due to scattering and providing an even field of radiant heat over the bed surface. The observation light provides intense light for procedures. The alarm light is located on the end of the heater assembly for ease of viewing. The entire heater assembly rotates to the side for X-ray procedures and for observation lamp replacement.

1.4 Control Unit

The control unit contains the electronic circuits and controls used to operate the radiant heater and the observation light. The control unit performs regular self checks during its operation including failure diagnostics.

Either manual or servo mode of operation may be selected. In the manual mode of operation, select the level of radiant heat output as indicated by the % power display on the control panel. The control circuit then maintains the selected level of radiant heat. The manual mode has a preheat setting which allows the Warmer to be pre-heated. In the servo mode of operation, select the patient's control temperature. A skin temperature probe is used to monitor the patient skin temperature. The control system modulates the radiant heat to maintain the patient at the selected control temperature. The patient's skin temperature is continuously displayed.

Alarms activate to alert the operator of a low or high patient temperature, a skin temperature probe failure, a power failure, equipment failure or a check patient prompt.

1.5 Bed Platform

Warmers with beds include a mattress and transparent side panels. The side panels fold down for easy access to the patient and can be removed for cleaning. The X-ray cassette tray (located in an opening beneath the bed when installed) pulls out for insertion of X-ray cassettes and allows X-rays to be taken without moving the patient.* Markings located along the side panels allow easy location of the cassette in the X-ray tray relative to the position of the patient on the warmer bed.* The tiltable bed platform allows Trendelenburg and reverse Trendelenburg positioning. The hydraulic system for the tiltable bed provides a smooth, dampened motion to avoid disturbing the patient.* A chest drainage hanger attached to the bed and rear cross bar extrusion provides a convenient location to hang chest drainage devices.*

1.6 Bassinet (3500 Series)

The bassinet unit can be detached and rolled away from the model 3500 warmer unit by pulling up on the locking lever knob located at the top rear of the bassinet (see Figure 1-1). The bassinet includes:

1. Transparent locking side panels which fold down for easy patient access and are removable for cleaning.

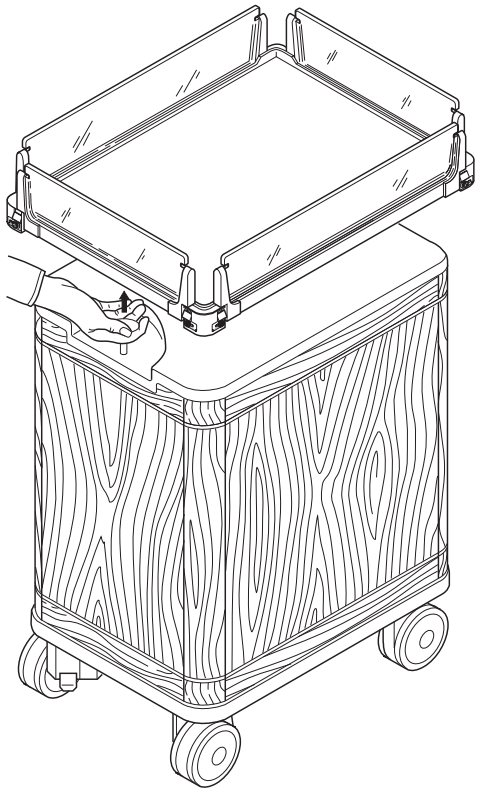
* Not available on the model 3500 warmer.

2. A mattress.

3. A tiltable, positive-lock bed platform which allows Trendelenburg and reverse Trendelenburg positioning at 4° and 8° tilt positions .

4. Three drawers which open from the front (or from the side in an optional configuration) for storing equipment or supplies. The top drawer is provided with a key lock.

5. Front locking casters.



CI.10.017

Figure 1-1
Operating the bassinet locking lever knob

1.7 Accessories

The rail mounting system is the basis for most of the accessories for the Warmer. This patented design consists of a dovetail shaped aluminum extrusion and a positive locking mounting block. Mounting blocks are attached to various accessories for mounting to the rail system. The mounting blocks are produced in two standard lengths and are machined to accommodate each specific accessory.

Accessories which mount on a single upright are interchangeable between all model Warmers.

Accessories can only be mounted to the inside dovetail rail on each upright of the 3500 Warmer.

Notes

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E. Interlock Switch Check 2-7

F. Power Failure, Memory and Battery Test (all models except the 2001 International) 2-7

2.1 Setup

Refer to the setup instructions shipped with the warmer for initial unpacking and setup of the unit after shipment. For model 3050, 3100 and 3150 wall mounted units, refer to the mounting instructions in the Appendix.

After removal from the shipping containers, inspect the Ohmeda Infant Warmer System and all accessory items for any signs of damage which may have occurred during shipment. File a damage claim with the shipping carrier if damage has occurred. Also confirm the presence of all accessory items or factory installed options as listed on the packing slip.

2.2 Mechanical Checkout Procedure

WARNING:

⚠ **Before using the Ohmeda Infant Warmer System, read this entire manual. Attempting to use this device without a thorough understanding of its operation may result in patient or user injury.**

⚠ **Do not perform the Checkout Procedures (Mechanical and Control Unit) while a patient occupies the warmer.**

⚠ **Complete the "Checkout Procedures" section of this manual before putting the unit into operation. If the warmer fails any portion of the checkout procedures it must be removed from use and repaired.**

A. Overall Appearance

1. Disconnect the power cord from the ac power source for the mechanical checks portion of this procedure.

2. Check the overall appearance of the Infant Warmer/ Bassinet System. There should be no obvious damage.

3. For the model 3500, separate the warmer from the bassinet.
4. For units with casters, check that all casters are in firm contact with the floor and that the warmer is stable and moves freely.

   For the model 3500, check that all six casters on the warmer move freely.

   Note: On the model 3500, it is possible that the two center casters may not be in contact with the floor at all times.
5. Lock the two front casters and check that the warmer is held in place. On the model 3500, lock the two rear casters and check that the warmer is held in place.
6. For the model 3500, place the bassinet on a level surface. Check that all four casters are in firm contact with the floor and that the Bassinet moves freely.
7. For the model 3500, lock the two front casters and check that the Bassinet is held in place.
8. Examine the power cord for damage. Replace the power cord if damage is evident.
9. Examine the unit for objects placed on top of the heater assembly.
10. Check heater-to-bed spacing falls within ranges specified in "1.3 Heater Assembly".

WARNING: ⚠ **Do not place any accessories or other objects directly over the bed surface. This may block radiant heat energy and lead to cooling of the infant.**

⚠ **Do not place items on top of the heater assembly. Items placed on top of the heater assembly can fall and injure the patient, prevent adequate ventilation of the heater assembly, and may pose a fire hazard.**

B. Heater Assembly Rotation

Rotate the Heater Assembly to the side and then back to the normal position. Check for a smooth rotation.

C. Mechanical Checks

1. Check the operation of the bed sides. The bed sides should operate smoothly.

WARNING: ⚠ **Regularly inspect the bed side panel latching mechanism, and the bedside locking mechanism on the model 3500, to ensure proper operation and to prevent the infant falling off the bed.**

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2. Check the operation of the tilt mechanism. Verify that the bed platform operates smoothly and locks in normal, Trendelenburg and reverse Trendelenburg positions.
3. If the drawer package is installed, check that all the drawers open and close freely. On units with the rotating drawer package, make sure the drawers rotate from front to 90° left and right and firmly stop in the detented positions.

WARNING: ⚠ **Overloading the drawers can affect the stability of the unit. Limit the load to 4.6 kg per drawer.**

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4. For the model 3500 bassinet, remove the keys from the top drawer and lock it. Check that the drawer is securely held closed.

D. Warmer/Bassinet Interlock (model 3500 only)

1. Lock the 3500 warmer's two rear casters.
2. Align the rear of the bassinet with the warmer's base.
3. Push the bassinet into the alignment/locking track of the warmer's base.
4. Continue pushing the bassinet until its locking pin drops into the socket at the end of the alignment track. There should be an audible click when the pin drops into position.
5. Lightly push and pull the bassinet to verify that the two units are securely held together.
6. Unlock the two rear casters on the warmer and check that the combined unit moves smoothly.

E. Warmer/Bassinet Unlock (model 3500 only)

1. Verify that the two bassinet front casters are unlocked and the two warmer rear casters are locked .
2. With the locking pin handle (located at the rear of the bassinet) raised, pull the bassinet forward until it is fully detached from the warmer .

CAUTION: ⚠ To prevent the drawers from opening unintentionally while moving the detached bassinet, move it from the front only.

F. Accessory Checks

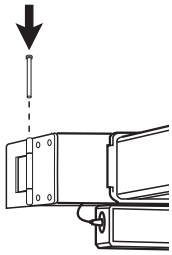
Perform these checks if they are applicable.

1. Check that all accessories are mounted securely and that the load limits are not exceeded.
2. Check that all gas accessories are installed and operating properly (refer to Section 3.9).
3. Where applicable, perform the checkout procedures detailed in the Operation and Maintenance Manuals for the accessories.

WARNING: ⚠ **Limit the load of accessories to 23 kg per side on the Warmer to ensure stability. Always try to evenly distribute the weight of accessories on both sides of the unit for a more balanced load. Accessories should not be mounted more than 142 cm above the floor. For models 3000 and 3500, limit the load of accessories to 9 kg maximum per side mounted no more than 112 cm above the floor.**

⚠ **Due to the increased height of units with the ECMO option installed, a tipping hazard may exist. Limit the total accessory load to 23 kg, no more than 11 kg per side.**

G. Wall Mount Checkout



CI.14.096

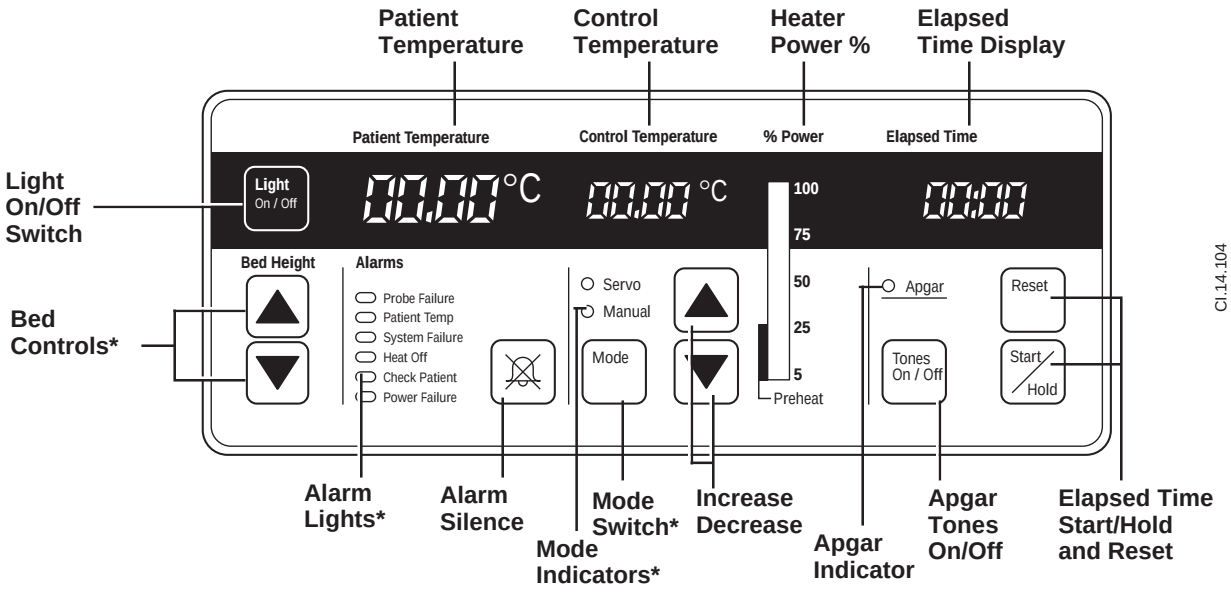
Figure 2-1
Wall mount pin insertion

1. Verify that the warmer is rigidly secured to the wall and the heater assembly is level.

   Note: To access the control unit and display module for service procedures, either hinge pin may be removed, allowing the warmer to pivot away from the wall.

2. Check that both hinge pins are in place and fully inserted with the pin heads at the top of the hinge.

WARNING: ⚠ **In the service position the strength of the hinge bracket is reduced. Never place a patient in the bed when the heater is in the service position. Never leave the unit unattended in the service position or with either pin removed.**



CI.14.104

* Note: Servo mode indicator, mode switch, and probe failure alarm are not present on the model 2001 international control panel. Bed raise/lower are only present on elevating models.

Figure 2-2
Control Panel

2.3 Control Unit Checkout Procedure

WARNING: ⚠ **Do not perform the Checkout Procedures (Mechanical and Control Unit) while a patient occupies the warmer.**

⚠ **Complete the "Checkout Procedures" section of this manual before putting the unit into operation. If the warmer fails any portion of the checkout procedures it must be removed from use and repaired.**

A. Control Unit Check

1. Connect the warmer power cord to an appropriate power source. Refer to the rating plate on the Warmer for the proper voltage needed. Switch the power On and verify the following on the Control Panel (Figure 2-1):
   a. The alternating two tone audible alarm sounds and all displays and indicators are lit for approximately two seconds.

      Note: During this time the controller also performs self check functions. If the controller detects a failure, the alarm stays on and service is required.

      Note: All alarms except system failure or power failure are preceded by a 30 second intermittent operator prompt tone.

   b. The manual mode indicator is lit (except model 2001).

   c. Operator prompt tones sound and the % power display flashes.

2. Adjust the heat output with the increase (▲) and decrease (▼) touch switches to attain the high and low limits as indicated by the % power display.

   Note: Steps 3 through 8 do not apply to the model 2001 International.

3. Connect the skin temperature probe to the warmer.

4. Press the mode touch switch to place the warmer in the servo mode and verify the following:
   a. The servo mode indicator is lit.

   b. An operator prompt tone sounds and the control temperature display flashes 36.5°C.

5. Press the increase (▲) touch switch and verify that the maximum servo control temperature attainable is 37.5°C.

   Note: A patient temperature alarm occurs if the difference between the patient temperature and the control temperature is greater than 1°C (the difference can be adjusted to 0.5°C by a qualified service person).

   Note: An alternating two tone alarm, a flashing overhead alarm light and a flashing patient temperature display may occur here if the skin temperature probe is below 30°C. Warm the probe with your fingers or silence the alarm.

6. Press the decrease touch switch and verify that the minimum servo control temperature attainable is 35.0°C.

7. Disconnect the skin temperature probe. Verify the following:
   a. The probe failure indicator light is lit.

   b. There is an alternating two tone alarm.

   c. The overhead alarm light is flashing.

   d. The patient temperature display flashes “HH.H”.

Note: The functionality of the remainder of the alarms is continually checked by the microprocessor software during normal operation. If a fault occurs in any of this circuitry, an indicator lights and a tone is sounded. The patient alarm can be verified manually by placing the unit in the manual mode and setting the power to 50%. After 12 minutes, the patient alarm should sound.

Additionally, a computer independent ”watchdog“ circuit will reset the computer, turn off the heater, and activate the alarms in the event of a microprocessor failure or software failure.

8. Press the alarm silence touch switch and verify the following:
   a. The probe failure indicator light is lit.
   b. The alternating two tone alarm is silenced.
   c. The overhead alarm light is lit.
   d. The patient temperature display indicates “HH.H”.
   e. After one minute, the alternating two tone alarm sounds, the overhead alarm flashes and the patient temperature display flashes “HH.H”.
9. Switch to the manual mode and set the heat at 25% power.

B. Elapsed Timer Check

1. Press the Start/Hold touch switch to activate the elapsed timer. Verify that the timer starts operating.
2. Press the Apgar tones On/Off touch switch for the Apgar tones. Verify that the indicator light for the Apgar tones is extinguished.
3. Press the Apgar tones On/Off switch for the Apgar tones again. Verify that the indicator light for the Apgar tones is lit.
4. Press the Start/Hold touch switch. Verify that the present elapsed time is held.
5. Press the Start/Hold touch switch and verify that the timer updates to the current elapsed time and the Apgar tones continue to sound at the specified times (at 1 minute and at every 5 minute interval after the elapsed timer has started).
6. Press the Reset touch switch and verify that the timer indicates “00:00”. If the elapsed timer is not used for approximately two minutes, the display switches off.

C. Observation Light Check

Press the Light On/Off touch switch. Verify that the observation light functions.

D. Raise and Lower Bed Switch Check

(Elevating Models Only)

CAUTION: ⚠ Do not continue to run the motor at the upper and lower limit positions; equipment damage may result.

---

1. Press the Raise Bed touch switch and verify that the bed raises to a maximum of 118 cm off the floor.
2. Press the Lower Bed touch switch and verify that the bed lowers to a minimum of 98 cm off the floor.

3. Press the foot switches on both sides of the unit and verify that they raise and lower the bed.

E. Interlock Switch Check

1. Place the warmer in the manual mode at 25% power output.

2. Rotate the heater assembly to the side. Verify that the heat off indicator light is On and the % power display indicates 0% heat.

3. Rotate the heater assembly to the normal operating position. Verify that the heat off indicator light is Off and the % power display indicates 25%.

F. Power Failure, Memory and Battery Test

1. Operate the unit in the manual mode with the heat set at 25% power for a minimum of one hour to charge the battery.

   Note: The battery must be fully charged to pass the 10 minutes test or partially charged to pass the two minute test. If the battery is defective, replace it. Refer to the service manual. There is no maintenance required for the battery. The battery has a two year replacement schedule.

2. Remove the Warmer power plug from the power source for two minutes. Do not switch the power Off. The power failure alarm should sound for two minutes and the power failure indicator on the display panel lights.

   Note: If the power failure alarm is tested for 10 minutes, the warmer must be connected to the correct power source and operated for 24 hours to recharge the battery before allowing a patient to occupy the Warmer.

   Note: The power failure alarm will not operate if the circuit breaker trips.

3. Reconnect the warmer to the power source. Verify the following:

   a. The warmer is operating in the servo mode (except model 2001).

   b. The heat output is still set at 25% power.

   c. The audio power failure alarm is off.

Notes

3/Operation

In this section

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3.1 Control Panel Operation

(Control Panel is shown in Figure 2-2)

A. Displays

- Patient Temperature
- Control Temperature
- % Power
- Elapsed Time

Patient Temperature:

The patient temperature display indicates the temperature sensed at the skin temperature probe tip. The skin temperature probe must be properly connected to the warmer and the infant for an accurate patient temperature measurement. The range of temperature measurement is from 30.0 to 42.0°C, with a resolution of 0.1°C. Temperatures above this range result in an “HH.H” patient temperature display and below this range result in an “LL.L” patient temperature display .

Control Temperature: (all models except the 2001 International)

The control temperature display indicates the control temperature you select in the servo mode. The servo control temperature range is from 35.0 to 37.5°C. The control temperature display is not active in the manual mode.

% Power:

The % power display indicates the percentage of maximum power that is being supplied to the radiant heater in 5% increments.

The preheat range (0 to 25% power) allows operation without the 12 minute check patient alarm in the manual mode.

Elapsed Time:

The elapsed time display indicates elapsed time in minutes and seconds up to a maximum of 60 minutes.

B. Indicator Lights

- Servo Mode Indicator
- Manual Mode Indicator
- Apgar Tones Indicator

Servo Mode Indicator: (all models except the 2001 International)

The servo mode indicator lights when the warmer is in the servo mode of operation. For proper operation of the warmer in the Servo Mode see section 3.2.

Manual Mode Indicator: (all models except the 2001 International)

The manual mode indicator lights when the warmer is in the manual mode of operation. For proper operation of the warmer in the Manual Mode see section 3.2.

WARNING: ⚠ **Use the servo mode unless the manual mode is specifically prescribed. While both modes require patient monitoring, the manual mode requires constant attention. In the manual mode, you must take the responsibility for making heater adjustments or environmental changes in response to the following conditions:**

- **patient size and weight;**
- **bed to heater head height differences;**
- **changes in the environment (drafts, direct sunlight, phototherapy lamp usage, etc.);**
- **changes in the patient condition**

In the servo mode, the warmer automatically adjusts heater output to maintain the desired skin temperature, reducing (but not eliminating) the need to monitor the patient and make environmental adjustments to ensure adequate patient temperature.

Apgar Tones Indicator:

The Apgar tones indicator lights when the Apgar tones are activated, and the Apgar timer is running.

C. Switches

- Mode Switch
- Increase (▲) and Decrease (▼) Switches

- Alarm Silence
- Start/Hold
- Reset
- Apgar Tones On/Off
- Light On/Off
- Raise bed (▲) (Elevating Models Only)
- Lower bed (▼) (Elevating Models Only)

Mode Switch: (all models except the 2001 international)

This switch is used to select either the Manual or Servo mode of operation. An audio tone sounds momentarily when the mode switch is depressed.

Increase (▲) and Decrease (▼) Switches:

These switches are used to set the radiant power level (% power) in the manual mode and to set the control temperature in the servo mode.

Alarm Silence:

This switch is used to silence the audible alarm. It silences all alarms except the system failure alarm. Refer to Table 1 - Warmer Alarms for details on how long the audible alarms are silenced.

Holding this switch depressed for five seconds or longer activates the audible alarm and lights all the indicator lamps for testing purposes.

Start/Hold:

This switch activates the elapsed timer. Refer to the elapsed timer operation, Section 3.3, for a detailed explanation of its function.

Reset:

This switch is used to reset the elapsed timer to “00:00”.

Apgar Tones On/Off:

This switch activates the Apgar tones. When the Apgar tones are activated, a short alarm tone is sounded at one minute and every five minute interval after the elapsed timer is started.

Light On/Off:

WARNING: ⚠ **Prolonged exposure to the light emitted by the observation lamp in this unit may harm the unprotected eyes of the infant. For safety, cover the infant’s eyes.**

This switch activates the observation light located in the heater assembly.

Raise Bed (▲) (Elevating Models Only)

WARNING: ⚠ **Check for proper clearance above the Warmer and below the bed surface before raising or lowering the bed.**

⚠ **Inspect all patient connected tubes or wires before and after moving or tilting the bed. Tilting or moving the warmer bed up or down can pull on tubing or leads connected to the patient. This may disconnect tubes or leads, restrict gas or liquid flow, or move sensors out of position.**

CAUTION: ⚠ Do not continue to run the motor at the upper and lower limit positions; equipment damage may result.

This switch raises the bed to a maximum height of 118 cm from the floor.

Lower Bed (▼) (Elevating Models Only)

This switch lowers the bed to a minimum height of 98 cm from the floor.

On/Off Power Switch and Circuit Breaker:

(See Figure 3-1.)

The On/Off power switch is located on the left side of the controller assembly (as viewed from the front) near the power cord socket. It is used for switching the power to the warmer(On and Off). The switch is also a circuit breaker and limits the maximum current drawn by the unit. If this circuit breaker trips when the warmer is operating, the power switch is deactivated to the Off position. To reset the circuit breaker, return the power switch to the On position. If the circuit breaker trips again, service is required.

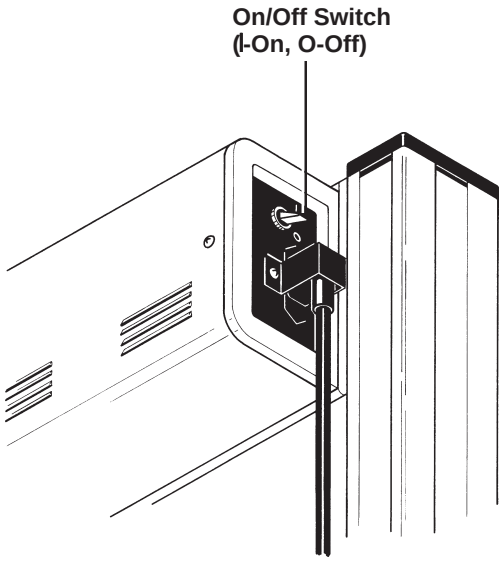

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Figure 3-1
On/Off Power Switch with built-in circuit breaker

D. Alarms

All alarms except system failure and power failure are preceded by a 30 second intermittent operator prompt tone.

- Probe Failure Alarm
- Patient Temperature Alarm
- System Failure Alarm
- Heat Off Alarm
- Check Patient Alarm
- Power Failure Alarm

Probe Failure Alarm: (all models except the 2001 international)

The probe failure alarm is only active in the servo mode of operation. The alarm activates when the skin temperature probe fails electrically due to an open or short circuit, or is disconnected from the Warmer. The heater deactivates and the patient temperature display flashes “HH.H” or “LL.L” when this alarm condition exists.

Patient Temperature Alarm: (all models except the 2001 international)

The patient temperature alarm is only active in the servo mode of operation. The alarm activates when the difference between the patient temperature and the control temperature is greater than 1°C. When the patient temperature returns to within 0.8°C of the control temperature, this alarm is automatically reset.

Note: If the patient temperature probe reads below 30°C or above 42°C in servo mode, the heater is deactivated and “LL.L” or “HH.H” respectively, appears in the patient temperature display.

Note: Qualified service personnel can adjust the alarm to trigger at a difference of 0.5°C and reset at a difference of 0.3°C.

WARNING:

⚠ **In the servo mode, verify that the patient temperature probe is securely attached to the patient at least once every half hour. A dislodged probe may not trigger an alarm. If the probe becomes dislodged, the warmer can over or under heat the infant.**

⚠ **The skin temperature probe should be located on the patient’s skin in an area which is directly in the path of the radiant heat. It should not be attached to an area which is shielded from the radiant heat or between the patient and the mattress. Large temperature gradients and very long servo response times will result from improper probe placement.**

⚠ **Rectal temperatures must never be used to servo control a patient’s temperature.**

System Failure Alarm:

The system failure alarm activates if the solid state relay controlling the radiant heater fails, if the microprocessor fails or if the calibration drifts by more than 0.3°C. The alternating two tone alarm cannot be silenced with the alarm silence switch.

WARNING:

⚠ **Do not use the Warmer system if the system failure alarm is activated. Remove the unit from service and refer to qualified personnel for repair.**

Heat Off Alarm:

When the heater is rotated to the side position, the heater is switched off and the Heat Off indicator is switched on. The audible Heat Off alarm activates after 5 minutes if the heater is not returned to the normal position. The alarm can be silenced by pressing the Alarm Silence switch.

Check Patient Alarm:

1. **Manual Mode:** The check patient alarm activates in the manual mode of operation when the heater has been energized at a power level greater than 25% for 12 minutes. The alternating single tone alarm is activated for this condition. Pressing the alarm silence switch silences this alarm and resets the timer for another 12 minutes of operation. If this alarm is not silenced within 3 minutes of the check patient alarm occurring, an alternating two tone alarm is sounded and the heater is deactivated until the alarm silence switch is pressed.

2. **Servo Mode:** The check patient alarm activates in the servo mode of operation if the heater has been at the 100% power level for 12 continuous minutes. The alternating single tone alarm is activated for this alarm. Pressing the alarm silence switch silences this alarm and resets the timer for another 12 minutes of operation. If this alarm is not silenced within 3 minutes of the check patient alarm occurring, the alternating two tone alarm is sounded and the heater is deactivated until the alarm silence switch is pressed.

Power Failure Alarm:

A battery operated power failure alarm activates when the external power source fails or is accidentally disconnected. The battery also provides power to an electronic memory to recall previous control temperature settings for approximately 10 minutes, when the battery is fully charged.

Table 1 — Warmer Alarms

Alarm Condition	Alarm Sound	Alarm Silence Period	Heater
Probe Failure‡	Alternating Two-Tone †	1 minute	Off
Patient Temperature is greater than 42°C‡	Alternating Two-Tone †	1 minute	Off
Patient Temperature more than 2°C from Control temperature‡	Alternating Single-Tone †	5 minutes	*
Patient Temperature between 1 and 2°C §of Control temperature‡	Alternating Single Tone †	15 minutes	*
Patient Temperature is less than 30°C ‡	Alternating Two-Tone †	1 minute	Off
System Failure**	Alternating Two-Tone	Cannot be silenced	Off
Heat Off	Alternating Single Tone †	5 minutes	Off
Check Patient	Alternating Single Tone † after 12 minutes	12 minutes	*
	Alternating Two-Tone after 15 minutes	15 minutes	Off
Power Failure	Alternating Single Tone	Cannot be silenced	Off

* Heater output is dependent on the Patient Temperature and Control Temperature settings in the servo mode, and the % Power setting in the manual mode.
** Error code in elapsed time display indicates the cause of the alarm.
† Preceded by a 30 second operator prompt tone.
‡ Not applicable to model 2001 International
§ Can be set to between .5 and 2°C by qualified service personnel.

Error Codes:

In the event of a system failure, a nonsilenceable, high priority alarm sounds and the System failure LED illuminates. Before shutting off the Warmer and removing it from service, make note of the error code that appears in the elapsed time display. This code will aid qualified service personnel in diagnosing the problem. Table 2 lists the possible error codes, for a more detailed explanation of error codes see the Service manual.

Table 2— Warmer Error Codes

E001	Instruction Test Failure	E002	ADC High Calibration Failure
E003	ADC Low Calibration Failure	E004	Checksum Failure
E005	RAM Test Failure	E007	ADC Converter Failure
E010	Line Voltage Out of Range	E012	Heater Not Switching On
E013	Heater Not Switching Off	E014	Alarm Oscillator Failure
E015	Software Upset - RAM Inconsistency	E016	Safety Relay Not Opening Properly
E017	Software Upset - Missing Routine	E019	Software Upset
E025	Hardware Watchdog Circuit Tripped		

3.2 Manual and Servo Mode Operation

WARNING:

⚠ **Before using the Ohmeda Infant Warmer System, read this entire manual. Attempting to use this device without a thorough understanding of its operation may result in patient or user injury. This device should only be operated by personnel trained in its operation and under the direction of qualified medical personnel familiar with the benefits and risks of this type of device.**

⚠ **Do not leave the patient unattended when using the Infant Warmer. Check the patient's temperature periodically using an independent temperature monitoring device to ensure the comfort and the safety of the patient. If the warmer is used for an extended time, it is recommended that the servo mode of operation be used. When an alarm is silenced, close monitoring of the patient's condition is required.**

⚠ **Use of electrosurgical units or other electrical field radiating equipment can affect the operation of the Warmer. Keep the patient probe lead as far away as possible from electrosurgical cables. Do not allow excess electrical cables to be laid on the bed platform. Use of electrosurgical units or other instruments which radiate electrical fields can cause indirect heating, by several tenths of a degree of the skin temperature probe due to absorbed electrical energy. When using these devices near the radiant warmer, operate the Warmer in manual mode for maximum safety.**

⚠ **The use of phototherapy equipment may raise the patient's temperature.**

⚠ **Radiant warmers increase an infant's insensible water loss. Take appropriate measures to maintain the patient's fluid balance while caring for them in a radiant warmer.**

WARNING:

⚠ **Do not use the Warmer in the presence of flammable anesthetics; a possible explosion hazard exists under these conditions.**

⚠ **Radiant energy can adversely affect blood components. When using intravenous tubing systems for delivery of blood components to patients occupying a warmer, shield any tubing with aluminum foil.**

⚠ **When using a radiant warmer, change the patient's diapers frequently. Radiant energy causes more rapid urine evaporation, and may lead to inaccurate urine diagnosis test analysis and inaccurate weight measurements.**

⚠ **Do not touch the protective grill under the radiant heater or the top of the heater assembly. These surfaces may be hot and a burn could result.**

⚠ **Always set the caster brakes before placing a patient in the warmer.**

---

1. Connect the power cord to a properly grounded AC power source.

2. Place the power switch in the On position. During the first seconds of operation, the warmer performs a self check of the control system. The software is verified, calibration is checked and operation of the solid state relay controlling the heater is verified. All displays and indicators are lit and the audible alarm is sounded. If the self check detects a failure, the alarm stays on and service is required.

   Note: The Warmer begins operation in the Manual Mode, described in Step 3. To operate the warmer in the Servo Mode, refer to the Servo Mode Operation in Step 5.

3. Manual Mode Operation.

WARNING:

⚠ **Use the servo mode unless the manual mode is specifically prescribed. While both modes require patient monitoring, the manual mode requires constant attention. In the manual mode, you must take the responsibility for making heater adjustments or environmental changes in response to the following conditions:**

- **patient size and weight;**
- **bed to heater head height differences;**
- **changes in the environment (drafts, direct sunlight, phototherapy lamp usage, etc.);**
- **changes in the patient condition**

In the servo mode, the warmer automatically adjusts heater output to maintain the desired skin temperature, reducing (but not eliminating) the need to monitor the patient and make environmental adjustments to ensure adequate patient temperature.

In the manual mode of operation, an operator prompt tone and the flashing of the % power display prompt you to select a level of radiant heat.

Use the increase (▲) or decrease (▼) touch switch to adjust the % power in 5% increments. The % power display indicates the power level selected. You must select a % power level each time the warmer is switched to the manual mode.

When you select % power levels between 30% and 100%, a check patient alarm sounds after 12 minutes to remind you to monitor the patient temperature. After checking the patient, this alarm can be reset by pressing the alarm silence switch.

This alarm recurs every 12 minutes during operation in the manual mode. If the alarm silence switch is not pressed within 3 minutes of the check patient alarm, an alternating two tone alarm sounds and the heater is turned off.

The Warmer may be preheated using 0% to 25% power levels. The 12 minute check patient alarm is not activated within these power level settings.

The skin temperature probe may be used to monitor the patient's temperature in the manual mode but it does not control the radiant heat energy level. See step 4 for skin temperature probe attachment.

Note: A patient placed in any warmer will normally develop temperature gradients with hotter and cooler areas. This is due to radiant heat being applied above the infant, the cooling effect from the mattress below the infant, the unequal skin cooling effect from evaporative water loss, unequal internal heat generation within the patient, and the environmental variables of room temperature, room air movement, incidental sunlight, etc.

4. Skin Temperature Probe Attachment.

WARNING: ⚠ **Use only the Reusable Ohmeda skin temperature probe (Stock No. 0208-0697-700) and heat reflective patches (Stock No. 0203-1980-300, 50/pkg) or the disposable probe (Stock No. 6600-0208-700, 10/pkg; Stock No. 6600-0196-700. 50/pkg) to monitor the patient's skin temperature. Use of other manufacturer's probes may affect the accuracy of warmer operation and the electrical safety of the patient.**

⚠ **The skin temperature probe should be located on the patient's skin in an area which is directly in the path of the radiant heat. It should not be attached to an area which is shielded from the radiant heat or between the patient and the mattress. Large temperature gradients and very long servo response times will result from improper probe placement.**

⚠ **Rectal temperatures must never be used to servo control a patient's temperature.**

The Ohmeda Patient probe lead is made from low mass wire that helps prevent probe detachment while reducing pulling on the neonate's skin.

Place the metal side of the skin temperature probe on the skin over the liver area of the infant's abdomen. Remove the paper protecting the hypoallergenic adhesive on the Heat Reflective Patch. Secure the skin temperature probe to the patient's skin with the adhesive side of the patch (Figure 3-2). Do not remove the heat reflecting foil which must be facing up.

Note: The probe jack is attached at a specific torque value. Loosening or tightening the jack may break the electrical connector. See Figure 3-3.

If the patient is prone, place the skin temperature probe on the back, where it will not be against the mattress. If the probe is between the patient and the mattress, it will produce false readings.

WARNING: ⚠ **Intimate contact between the skin temperature probe tip and the patient's skin must be maintained for accurate skin temperature measurement. Underheating or overheating may result from poor contact between the skin temperature probe and the patient. Verify that the skin temperature probe is securely attached to the patient at least once every half hour.**

Connect the skin temperature probe to the Warmer by plugging the probe connector into the left side of the display module as viewed from the front. (See Figure 3-3.)

CAUTION: ⚠ Always remove the probe from the patient by grasping and removing the heat reflective patch first, then remove the probe from the patient or the patch. Always remove the probe from the Warmer by grasping the plug at the panel. Placing excessive strain on the skin temperature probe lead wire can damage the probe.

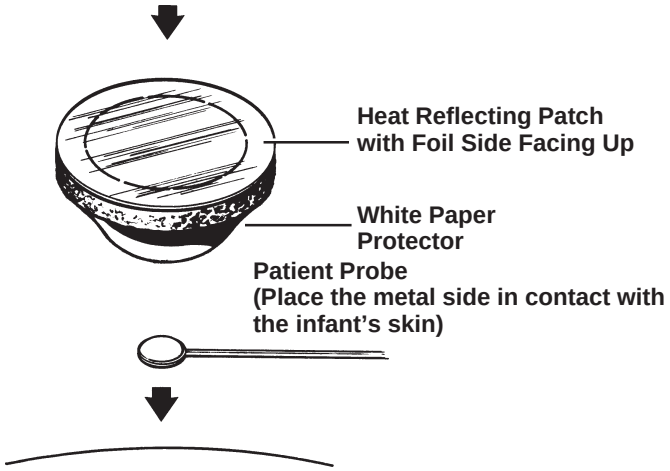

Figure 3-2
Skin Temperature Probe attachment

Note: Avoid placing excessive strain on the skin temperature probe lead. Always remove the probe by grasping the plug at the panel. Do not pull on the probe lead wire.

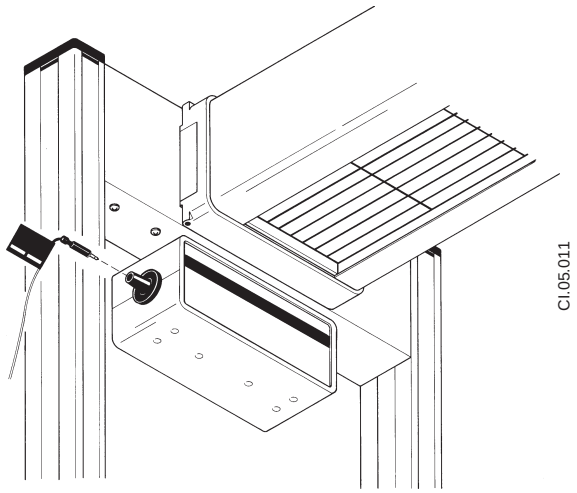

Figure 3-3
Temperature Probe connection to the Display Module

5. Servo Mode Operation. (all models except the 2001 International)

   Note: The skin temperature probe must be properly attached before starting servo mode operation. Refer to step 4.

   You must select a servo control temperature setting when the Warmer is used in the servo mode for the first time. You are prompted to make this setting with an operator prompt tone and the flashing of the control temperature display.

The servo control temperature is adjusted by pressing the increase (▲) or the decrease (▼) touch switches. The control temperature can be adjusted from 35 to 37.5°C. In the servo mode, the temperature sensed by the skin temperature probe is used by the control system to modulate the radiant heat and maintain the patient's skin temperature at the selected control temperature. The Appendix details the Servo Control Mode Algorithm.

Note: The Warmer cannot differentiate between an increase in core temperature with cold skin (fever), and low core and skin temperature (hypo-thermia). Patient temperature should be verified with an axillary thermometer.

WARNING: ⚠ **In the servo mode, verify that the patient temperature probe is securely attached to the patient at least every half hour. A dislodged probe does not always trigger an alarm. If the probe becomes dislodged, the warmer can over or under heat the infant.**

Note: A patient placed in any warmer will normally develop temperature gradients with hotter and cooler areas. This is due to radiant heat being applied above the infant, the unequal skin cooling effect from evaporative water loss, unequal heat generation within the patient, and the environmental variables of room temperature, room air movement, incidental sunlight, etc.

WARNING: ⚠ **The use of phototherapy equipment may raise the patient's temperature.**

3.3 Elapsed Timer Operation

1. Press the Start/Hold touch switch to activate the elapsed timer.
2. The Apgar indicator light is illuminated when the Apgar tones are enabled. Press the Apgar Tones On/Off touch switch to enable or cancel the Apgar tones.
3. Press the Start/Hold touch switch to hold the present elapsed time when the timer is running. The Apgar tones, if enabled, continue to sound at the specified times (after 1 minute and at every 5 minute interval after the elapsed timer is started).
4. Press the Start/Hold touch switch to update the timer to the current elapsed time.
5. Press the reset touch switch to start the timer over at "00:00". The elapsed time display is blanked out when the elapsed time reaches 60 minutes.

3.4 Bassinet Operation (model 3500 only)

Note: Lock the warmer's two rear casters before unlocking or locking the bassinet.

The bassinet unit can be detached from the model 3500 warmer by pulling up on the locking lever knob located at the top rear of the bassinet. When the knob is pulled up, the locking pin is freed from its socket on the warmer alignment/locking track on the warmer base. The bassinet can then be rolled forward away from the warmer.

To return the bassinet to its locked position, align the bassinet with the warmer . Push the bassinet over the warmer's alignment/locking track. Continue pushing the bassinet until its locking pin drops into the socket at the end of the alignment/locking track. There should be an audible click when the pin drops into position. Lightly push and pull the bassinet to verify that the two units are securely held together.

The bassinet's front casters can be locked in position by pressing the locking button on each caster.

3.5 Bed Platform Operation

(See Figure 3-4.)

On large and small bed units, the bed platform tilts for Trendelenburg and reverse Trendelenburg positioning capabilities. Lift or push down on the tilt handle to position the bed.

WARNING: ⚠ **Inspect all patient connected tubes or wires before and after moving or tilting the bed. Tilting or moving the warmer bed up or down can pull on tubing or leads connected to the patient. This may disconnect tubes or leads, restrict gas or liquid flow, or move sensors out of position.**

Note: Tilting the bed mattress can affect the operation and performance of the radiant warmer.

The model 3500 bassinet bed platform also tilts for Trendelenburg and reverse Trendelenburg positioning. To raise the front of the bed platform, pull the handle out and lift the bed. Release the handle so it engages in the upper detent. To lower the front of the bed platform, pull the handle out and lower the bed. Release the handle so it engages in the lower detent.

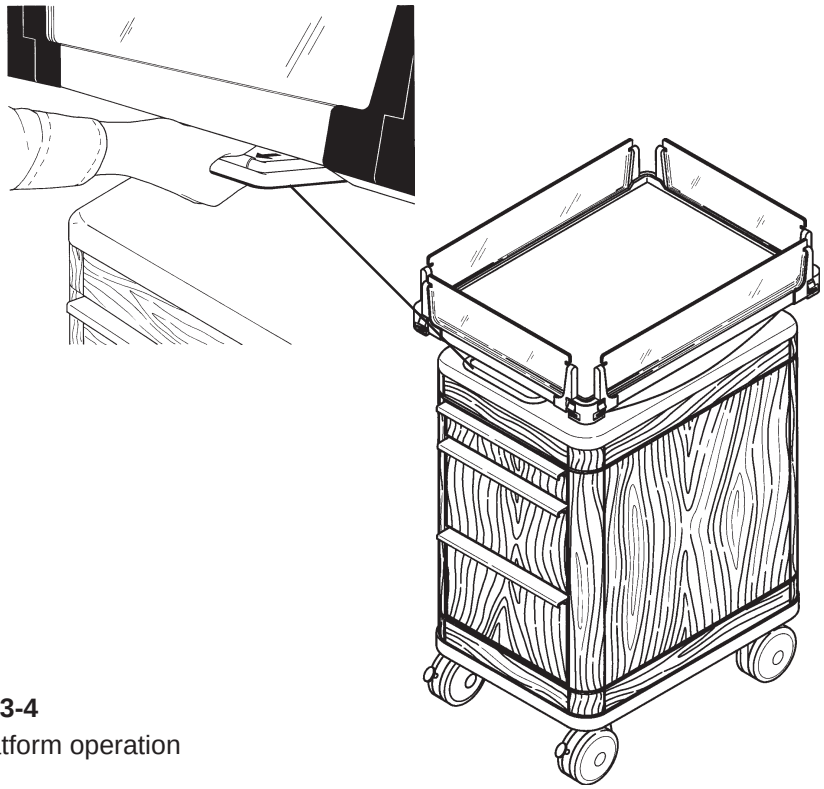

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Figure 3-4
Bed platform operation

3.6 Side Panel Operation

(See Figure 3-5.)

WARNING:

⚠ **Do not leave the patient unattended when the side panels are lowered.**

⚠ **Do not move the warmer by pushing or pulling on the bedside panels. This action may lead to the deterioration and breakage of the components which form a safety barrier around the infant.**

⚠ **Ensure that the bedside panels are locked in position when a patient occupies the bed. Blankets or other foreign objects may prevent the latches from fully engaging.**

To lower a side panel, pull it up and then pull the top edge away from the bed. On the model 3500 bassinet, the side panel locking mechanism must first be unlocked.

To raise a side panel, swing it to the upright position; then allow it to engage in the latched position.

To remove a side panel, lower the side panel, press the end pins in and lift the side panel out.

To replace a side panel, hold the end pins in, place the side panel in position and release the end pins.

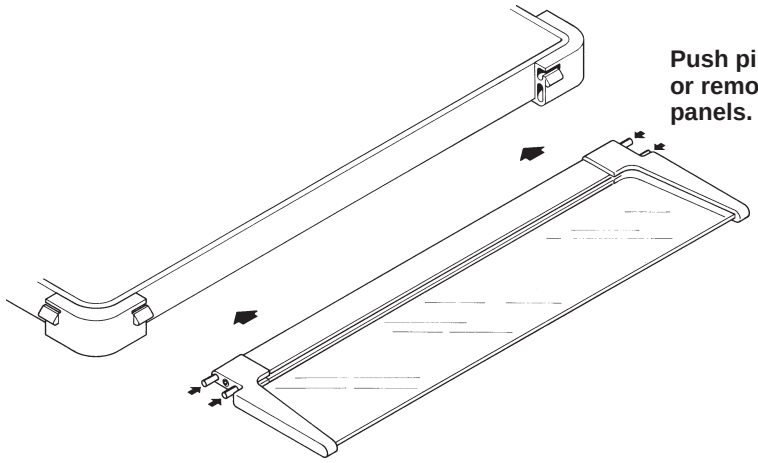

Figure 3-5
Side Panel operation

The Tubing Organizer Rear Side Panel aids in routing tubing to patients receiving therapy in warmer beds. The various size holes in the panel can accommodate large diameter tubes, such as aerosol tubes or the fiber optic cable of a BiliBlanket™ System; or smaller diameter tubes, such as ventilator or I.V. tubes. See Figure 3-6.

WARNING:

⚠ **Install tubing in the appropriately sized holes. Use of inappropriate holes may cause kinking, pinching or restriction of flow through the tubes, and may interfere with the proper operation of therapy equipment.**

⚠ **Do not install chest drainage tubes in the Tubing Organizer Rear Side Panel.**

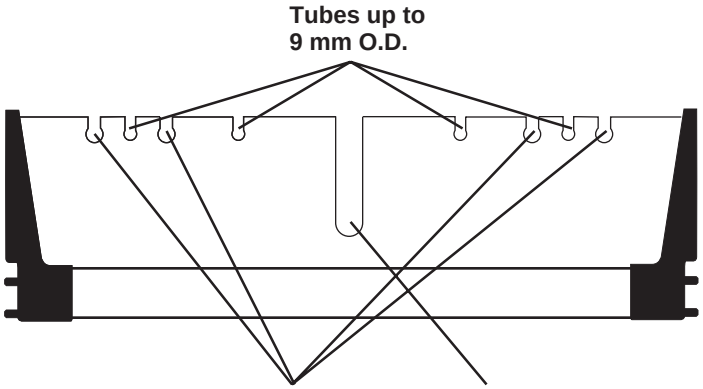

Figure 3-6
Tubing organizer panel

WARNING:

⚠ **Inspect all patient connected tubes or wires before and after moving or tilting the bed. Tilting or moving the warmer bed up or down can pull on tubing or leads connected to the patient. This may disconnect tubes or leads, restrict gas or liquid flow, or move sensors out of position.**

⚠ **Do not lower the Tubing Organizer Rear Side Panel with tubing attached. Lowering can pull on the tubing, causing the tubes to dislodge from the patient.**

⚠ **Radiant energy can adversely affect blood components. When using intravenous tubing systems for delivery of blood components to patients occupying a warmer, shield any tubing with aluminum foil.**

Using the side panel as a tubing organizer restricts the panel's ability to be lowered. Install the tubing organizer side panel on the back of the bed platform (the side nearest the aluminum uprights) to maintain easy access to the infant.

The tubing organizer side panel should be inspected, visually and by touch, for sharp edges, cracks or crazing, before each use.

3.7 X-ray Procedures

(See Figure 3-7 and 3-8)

The X-ray cassette may be placed in the slot under the bed platform or in the X-ray cassette tray if installed.

The heater housing rotates to the left for X-ray procedures. To place the heater housing in the X-ray position rotate the heater to the left. The heater is deactivated when in the X-ray position and the heat off indicator is illuminated. The audible alarm is activated after 5 minutes (in the manual mode) if the heater housing is not returned to the normal position. The alarm may be silenced for 5 minutes by pressing the alarm silence switch.

Note: The normal servo temperature alarms and alarm silence periods are in effect in the servo mode of operation. The 5 minute heat off alarm silence period is overridden by the normal servo temperature alarms. The heater is deactivated in the X-ray position. Rotate the heater housing to the normal position to resume normal operation.

The X-ray cassette tray (see Figure 3-8) used with all large and small bed warmers (except model 3500) facilitates X-ray procedures while patients occupy the warmer bed. An X-ray cassette can be placed on the tray and slid into the cavity beneath the bed without disturbing the patient. The tray can also be used as a writing surface or as work space during procedures, provided that a maximum load limit of 5 lbs is not exceeded.

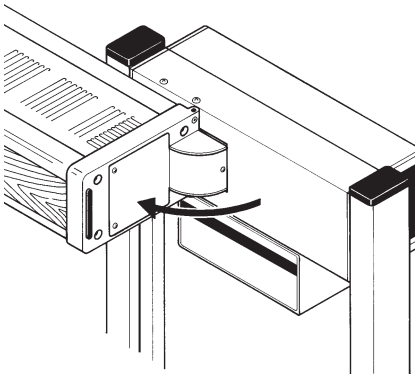

Figure 3-7
Heater Rotation for X-ray Procedures

Using the tray

WARNING: ⚠ **Limit the load placed on X-ray tray 2.3 Kg to avoid a tipping hazard.**

Never place an infant on the X-ray cassette tray.

1. To help position the X-ray cassette, the tray has detents or stops at its center position and at normal extension. Gently pull the tray out allowing gravity to engage the stops. Since the tray removes easily for cleaning, use caution when pulling it out, and properly support the tray and any X-ray cassette. The tray can be pulled out on either side of the warmer bed.

2. To further aid in positioning the X-ray cassette, a location grid pattern is molded into the tray itself. The tray grid markings correspond to the decal grid markings on the warmer bed side panels. Using the markings on the side panels as a reference to the infant's location, position the cassette on the tray so that, when the tray is fully inserted, the cassette will be directly beneath the infant.

3. With a smooth continuous motion, slide the tray back under the warmer bed until you feel the "click" when the stops on the slides engage the tray at the fully inserted center position.

4. Rotate the heater housing out of the way, position the X-ray machine and take the X-ray.

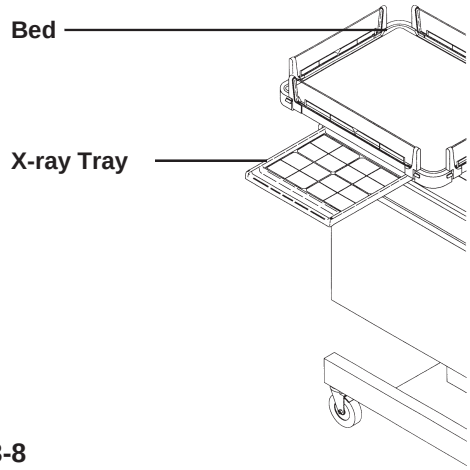

Figure 3-8
X-ray tray operation

WARNING: ⚠ **Do not place any foreign objects on the warmer bed or in the under bed cavity while performing X-ray procedures. Incompatible materials in the path of the X-ray may adversely affect the quality of the X-ray image. Use of mattress or bedding materials other than those supplied by Ohmeda should be evaluated by a Neonatologist or Radiologist.**

The tray can be removed for cleaning by simply sliding it all the way out of the slides. The tray should be cleaned between patients according to the hospital protocol . Refer to section 4.1 for a list of recommended cleaning agents.

3.8 Oxygen Administration

WARNING: ⚠ **Oxygen concentrations higher than 40% can increase the risk of retrolental fibroplasia (retinopathy of prematurity). It is probable that even concentrations of 40% or less oxygen (formerly considered safe) could be dangerous to some infants. Therefore, arterial blood gas measurements are extremely important for regulation of the concentration of inspired oxygen when an oxygen-enriched environment is considered necessary. (See current edition of "Standards and Recommendations for Hospital Care of Newborn Infants" prepared by the Committee of Fetus and Newborn of the American Academy of Pediatrics.)**

Oxygen can be administered from a regulated pipeline source or from auxiliary oxygen tanks mounted on the Warmer.

3.9 Gas Cylinder Installation and Operation

(See Figure 3-9)

Note: Use United States "E" size pin indexed gas cylinders only.

U.S. "E" size cylinders, with 11 cm x 66 cm outside dimensions, may be used with warmer units with elevating pedestals.

British "E" size cylinders are too long to be used with units with elevating pedestals. Mounting these larger cylinders on elevating units can damage the yoke, manifold or cylinder pin if the cylinder contacts the floor.

WARNING: ⚠ **Using medical gas cylinders that are not recommended for mounting on the Infant Warmer System may result in a fire hazard or an explosion.**

Note: Discontinue therapy while replacing cylinders.

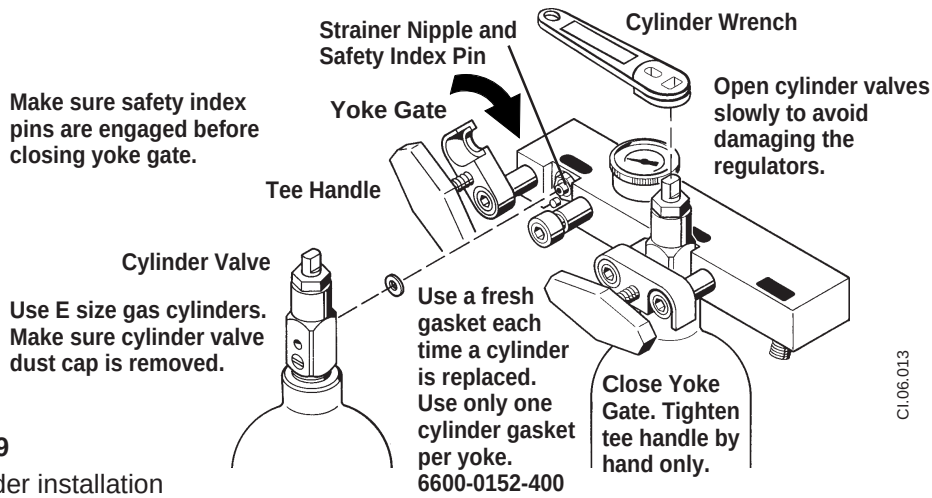
Strainer Nipple and Safety Index Pin
Cylinder Wrench
Make sure safety index pins are engaged before closing yoke gate.
Yoke Gate
Open cylinder valves slowly to avoid damaging the regulators.
Tee Handle
Cylinder Valve
Use E size gas cylinders. Make sure cylinder valve dust cap is removed.
Use a fresh gasket each time a cylinder is replaced. Use only one cylinder gasket per yoke. 6600-0152-400
Close Yoke Gate. Tighten tee handle by hand only.
CI.06.013

Figure 3-9
Gas cylinder installation

Cylinder Removal:

1. Close the valve on the cylinder to be removed.
2. While supporting the cylinder, back out the Tee handle until the tip of the screw is flush with the inside surface of the gate.
3. Swing the gate open in a counter clockwise direction.
4. Remove the old cylinder.
5. Remove the old gasket from the yoke strainer nipple and discard.

Cylinder Installation:

1. Install a new gasket (Ohmeda Stock No. 6600-0152-400) on the strainer nipple.

CAUTION: ⚠ Use only one cylinder gasket per yoke. Use of more than one gasket could cause leakage of the cylinder gas.

---

2. Before installing a cylinder:
   a. Remove the cylinder dust cap, if present.
   b. Briefly open then immediately close the cylinder valve to blow any foreign matter out of the valve outlet. Do not use excessive force on the shut off valve.
3. Install the cylinder valve over the strainer nipple, making sure that the safety index pins are engaged.
4. Swing the gate closed in a clockwise direction and tighten the Tee Handle sufficiently (by hand only) to hold the cylinder firmly in place. Do not use wrenches or any other tool on the Tee Handle screws.
5. If only one oxygen cylinder is used, a yoke plug (Ohmeda Stock No. 6600-0399-500) and a gasket are required to seal the unused yoke cylinder port.

CAUTION: ⚠ Yoke check valves are not intended to provide a leak free seal; always use a yoke plug and a fresh cylinder gasket to seal an unused cylinder port.

---

6. Open cylinder valve S-L-O-W-L-Y, and rotate until it is fully open.

CAUTION: ⚠ Open cylinder valves S-L-O-W-L-Y to avoid damaging the regulators.

Leak Testing the System:

Prior to initial use and as prescribed by hospital protocol, the yoke system(s) should be checked for leaks according to the following procedure:

l. Discontinue use of the system to be tested.

2. Disconnect all pipeline gas supply and outlet lines from the yoke assembly.

3. Use new gaskets and full cylinders at each cylinder port (optionally, one full cylinder and a yoke plug may be used on a 2 cylinder oxygen manifold)

4. Open a cylinder valve at each manifold to charge the manifold(s) to a minimum of 5137 kPa and then close the valve(s).

5. Note the gauge pressure reading in the manifold(s).

6. After one minute, check the gauge pressure reading(s) again. The pressure gauge needle should not drop visually. If there is a visual pressure drop, discontinue use of the gas system and have the yoke assembly repaired by a qualified service person.

Using Two Cylinders as an Oxygen Supply:

When two cylinders are installed to provide an oxygen supply, a check valve permits replacement of one depleted cylinder while the other continues to furnish oxygen. Leave the valve on the new cylinder closed until the old cylinder requires replacement, then open the valve on the new cylinder and immediately replace the old cylinder as described above. This assures that a full back-up cylinder is always available for use.

WARNING: ⚠ **Leaving both oxygen cylinder valves open at the same time will allow simultaneous depletion of both cylinders with no reserve oxygen supply available.**

Use of Pipeline Air and Oxygen Supplies:

1. Connect pipeline supply(s) to the appropriate pipeline inlet(s).

2. Connect desired equipment to the appropriate outlet(s).

3. Ensure correct operation of the secondary pipeline equipment.

When pipeline air or oxygen supplies are used, install cylinders as described above to provide an emergency gas supply in the event of a pipeline failure. Cylinder valves should remain closed until and unless the emergency gas supply is needed.

WARNING: ⚠ **Do not leave gas cylinder valves open if the pipeline supply is in use. Pressures from both oxygen supplies may become equal, and if simultaneously used, cylinder supplies may be depleted, leaving no reserve supply in case of pipeline failure.**

3.10 Mounting Accessories

(See Figure 3-10)

WARNING: ⚠ **Limit the load of the accessories to 23 kg per side on the Warmer to ensure stability. Always try to evenly distribute the weight of accessories on both sides of the unit for a more balanced load. Accessories should not be mounted more than 142 cm above the floor. For models 3000 and 3500, limit the load of accessories to 9 kg maximum per side mounted no more than 112 cm above the floor.**

⚠ **Due to the increased height of units with the ECMO option installed, a tipping hazard may exist. Limit the total accessory load to 23 kg, no more than 11 kg per side.**

Rail system components mount to the uprights and provide ready access to commonly used equipment such as suction regulators, flowmeters, collection bottles, etc.

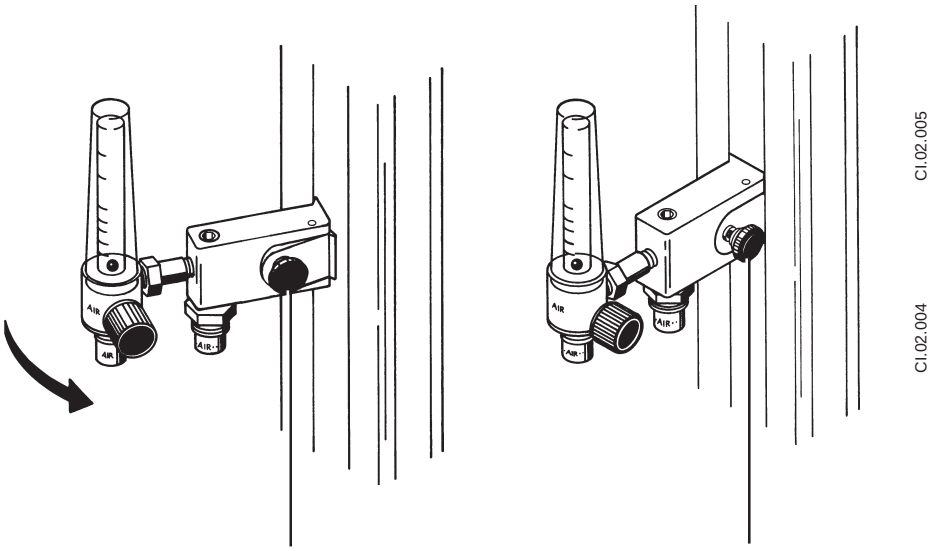

Figure 3-10
Mounting rail system components

A. Mounting and Releasing Procedure

(See Figure 3-10)

1. Loosen the mounting screw on the mounting block.
2. Place the mounting block in position on the rail.
3. Tighten the mounting screw.
4. Release the rail system component by loosening the mounting screw.

B. Adapter Plate Mounting and Releasing Procedure

(See Figure 3-11)

Universal Adapter plate allows Vacuum/Collection bottle slides and Bird blender bracket to mount to the rail system.

1. Loosen the two mounting screws on the side of the adapter plate with the hex key provided with the plate.
2. Place the adapter plate in position on the rail.
3. Tighten the two mounting screws.
4. Release the adapter plate by loosening the mounting screws.

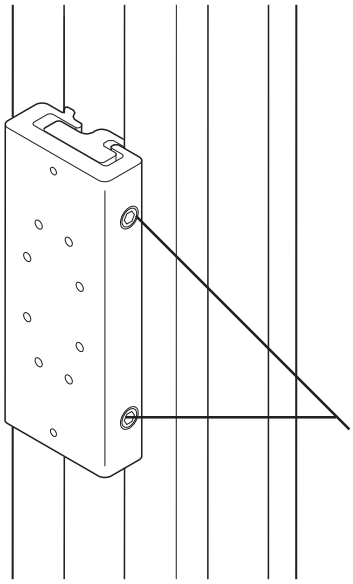

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Figure 3-11
Mounting the Adapter Plate

C. ECMO Adapter

The ECMO adapter raises the bed an additional six inches to facilitate Extra-Corporeal Membrane Oxygeneration procedures on model 4400 and 5000 units.

WARNING: ⚠ **Due to the increased height of units with the ECMO option installed, a tipping hazard may exist. Limit the total accessory load to 23 kg, no more than 11 kg per side.**

3.11 ThermaLink Option

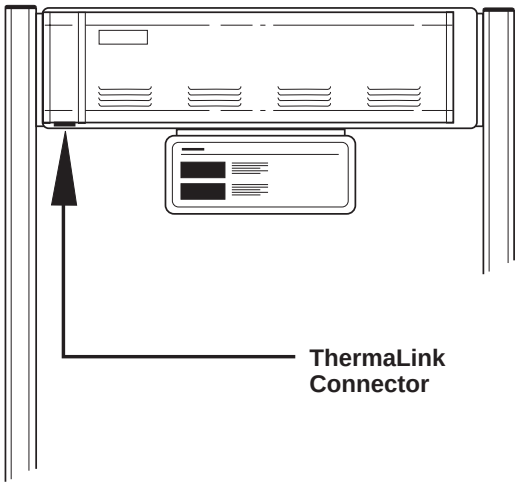

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Figure 3-12
ThermaLink connector (rear view of warmer)

The ThermaLink Serial data and Nurse Call connections are options offered with the Warmer (units with version 5.0 software or higher) . Your unit has this option if there is a nine pin connector on the back left side of the controller cover. See Figure 3-12.

Using the Serial Data interface

WARNING: ⚠ **Remote monitoring does not replace the need for direct patient observation by qualified medical personnel.**

The serial data output can be used with a computer or a commercial RS-232 monitor. Because of the wide variety of applications and systems, detailed information on decoding the data stream appears in the appendix. For details of the RS-232 protocol and connector pin out, refer to the Specification section.

WARNING: ⚠ **The computer or RS-232 monitor's user program must continuously check the data link. The program should constantly verify connection to the warmer controller and check for updated data.**

Using the Nurse Call System interface

WARNING: ⚠ **Remote monitoring does not replace the need for direct patient observation by qualified medical personnel.**

⚠ **If you connect the Nurse Call output to a system which uses a normally open connection, a disconnected Nurse Call cable will not trigger an alarm.**

The Nurse Call connector lets you use the Warmer with your current remote alarm system. Nurse Call alarms trigger for:

- Probe Failure Alarms
- Patient Temperature Alarms
- System Failure Alarms
- Heat Off Alarms
- Check Patient Alarms
- Power Failure Alarms

The Nurse Call alarms work with the warmer audible alarm. Silencing the audible alarm on the warmer stops the Nurse Call alarm even if the alarm condition still exists. At the end of the silence period, the Nurse Call alarm and the audible alarm reactivate unless the condition has been resolved. The alarm silence period ends prematurely if another alarm triggers.

Refer to the Appendix for additional information on Nurse Call connections.

Nurse Call checkout

1. Complete the checkout procedure in section 2.3.
2. Verify proper operation of the Nurse Call station.
3. Connect the Nurse Call connector to the warmer.
4. Unplug the warmer to trigger an alarm. Verify that you also get an alarm at the Nurse Call station.

Note: Any interruption of warmer power (deliberately switching off the warmer, accidently unplugging the power cord, etc.) triggers a Nurse Call alarm.

3.12 Rotating Drawer Option

The rotating drawer option allows access to the storage drawers from the front or from either side of the warmer. The drawer cabinet rotates by pushing on the side of the cabinet, stoping at 90° detent positions.

WARNING: ⚠ **Use caution when rotating the cabinet to avoid damage to the drawers or possible injury. Always ensure the drawers are fully closed before rotating the cabinet.**

Note: The rotating drawer option is not available on model 3400 or 3500 warmers.

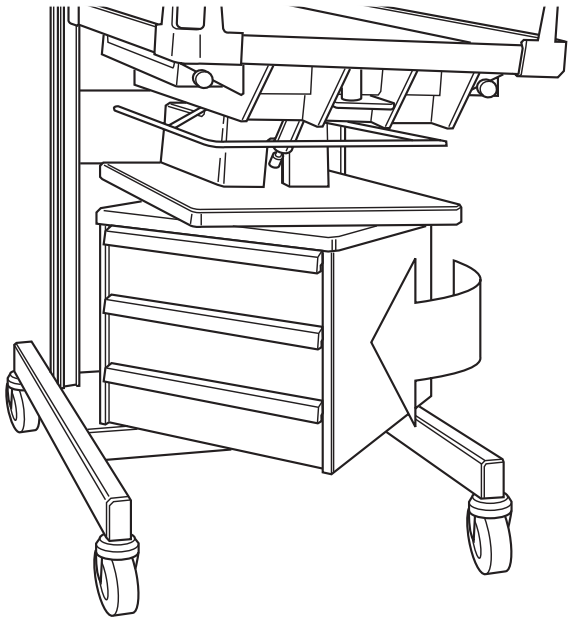

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Figure 3-13
Rotating Drawer Package

4/Cleaning and Disinfecting

In this section

4.1 Cleaning 4-1

4.2 Wood Surfaces (model 3500 only) 4-2

4.3 Reusable Skin Temperature Probe 4-2

4.1 Cleaning

WARNING: ⚠ **Disconnect power to the Warmer and allow the heat rod to cool before cleaning to avoid the possibility of a burn.**

Clean the Warmer at least once a week or after each patient. Cleaning procedures for the Warmer and accessories are explained in the following sections.

CAUTION: ⚠ Use the cleaning solution sparingly on a cloth when cleaning the warmer. Do not saturate the unit - excessive solution causes damage to internal components.

1. Unplug the power cord.

2. Clean the exterior of the warmer using mild detergent solution applied with a damp cloth or sponge. Aqueous solutions which are U.S. Environmental Protection Agency approved hospital level disinfectants may be used.

   Do not allow liquids to seep into the electrical housing.

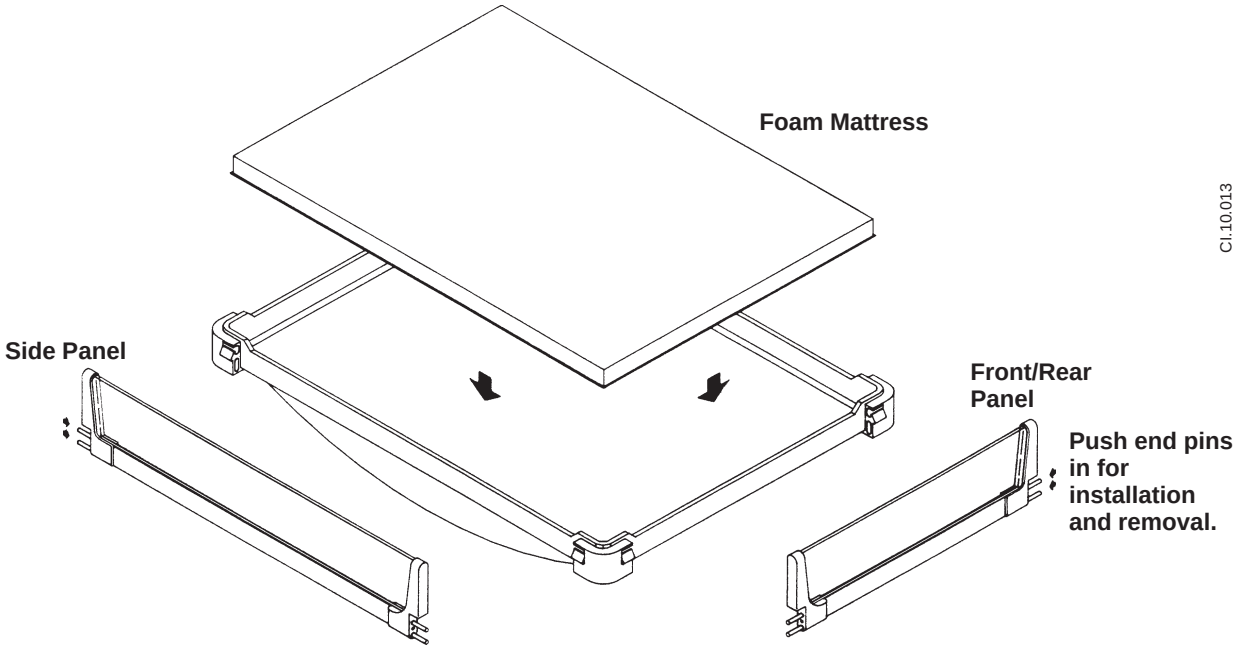

Figure 4-1
Disassembly for cleaning

3. The mattress, X-ray tray, bed, and side panels may be cleaned without immersing by using a disinfecting agent safe for use on the materials.

   Note: Bleach (sodium hypochlorite) should not be used tthe soft heater end cap.

   The following lists recommended cleaning solutions that may be used safely.

Generic Formulation	**Max. concentration level**
Hydrogen peroxide	6 %
Sodium hypochlorite compound	0.5 % Aqueous Solution
Quaternary ammonium	0.2 %
Iodophors	50 mg/liter
Glutaraldehyde	2 %

CAUTION:

⚠ Use of cleaning/disinfecting solutions containing chemicals that are not listed above (i.e. alcohol, acetone, etc.), or chemicals in greater concentrations than those listed above, may damage the patient probe or other material being cleaned.

⚠ Do not autoclave or gas sterilize the mattress.

4.2 Wood Surfaces

The wood surfaces are covered with a tough polymer coating which does not require waxing or special cleaners. Clean all wood surfaces with a mild soap and warm water solution.

Apply the cleaning solution with a clean cloth or sponge.

Dry the wood surfaces with a clean, soft cloth.

Dust the wood surfaces with a clean, soft cloth which is free of abrasive material.

4.3 Reusable Skin Temperature Probe

Clean the skin temperature probe by gently wiping with a soft, damp cloth containing detergent or disinfecting solution.

The table in section 4.1 (above) lists recommended cleaning solutions that may be used safely.

CAUTION:

⚠ Do not autoclave or gas sterilize the skin temperature probe. Do not immerse the probe in liquid cleaner. Avoid placing excessive strain on the probe lead. Always remove the probe by grasping the plug at the panel. Do not pull on the probe lead. These precautions will help avoid damage to the probe.

Note: Some cleansers may discolor the skin temperature probe over time.

Note: Disposable skin temperature probes cannot be cleaned.

5/Maintenance

In this section

5.1 Repair Policy and Procedure 5-1

5.2 Circuit Breaker Reset 5-2

5.3 Lamp Replacement 5-2

A. Alarm Lamp Replacement 5-2

B. Observation Lamp Replacement 5-3

5.4 Yoke Manifold Assembly 5-5

5.5 Battery Test 5-5

5.6 Maintenance Schedule 5-6

A. Operator Maintenance 5-6

B. Service Maintenance 5-6

5.1 Repair Policy and Procedure

Warranty repair and service must be performed by an Ohmeda Medical Service Representative or at the Datex-Ohmeda Service and Distribution Center (USA). To contact an Ohmeda Medical Service Representative, contact the nearest Ohmeda Medical Service Office.

Do not use malfunctioning equipment. Make all necessary repairs or have the equipment repaired by an Ohmeda Medical Service Representative. Parts listed in the Service Manual, for this product, may be repaired or replaced by a competent, trained person who has experience in repairing devices of this nature. After repair, test the equipment to ascertain that it complies with the published specifications.

CAUTION:

⚠ Only competent individuals trained in the repair of this equipment should attempt to service it as detailed in the Service Manual (Stock Number 6600-0195-000).

⚠ Detailed information for more extensive repairs is included in the service manual solely for use by individuals having proper knowledge, tools and test equipment, and for service representatives trained by Datex-Ohmeda.

Contact the nearest Ohmeda Medical Regional Service Office for assistance or before returning equipment for servicing. Obtain a Return Authorization Number before retuning equipment.

If you send equipment to the Datex-Ohmeda Service and Distribution Center, package it securely for protection, in the original shipping container if possible, and ship it prepaid. Enclose the following five items:

1. A letter describing in detail any difficulties experienced with the equipment. Reference the Return Authorization number
2. Warranty information — a copy of the invoice or other applicable documentation must be included.
3. Purchase order number to cover repair of equipment not under warranty.
4. Ship to and bill to information.
5. Person (name and telephone number) to contact for functional questions.

In all cases, other than where Ohmeda Medical's warranty is applicable, repairs will be made at Ohmeda Medical's current list price for replacement part(s) plus a reasonable labor charge.

5.2 Circuit Breaker Reset

(See Figure 5-1.)

The warmer is equipped with a combination power switch and manual-reset circuit breaker located on the left side of the controller assembly (as viewed from the front), near the power cord socket. The circuit breaker limits the maximum current drawn.

If this circuit breaker trips when the warmer is operating, the power switch is deactivated to the Off position. To reset the circuit breaker, return the power switch to the On position. If the circuit breaker trips again, service is required.

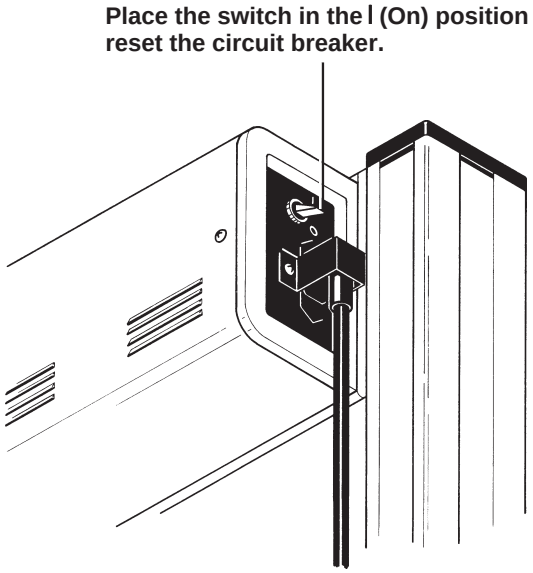

Figure 5-1
Circuit breaker reset

5.3 Lamp Replacement

WARNING: ⚠ **Disconnect the Warmer power cord and allow the unit to cool before replacing the alarm or observation lights.**

A. Alarm Lamp Replacement

(See Figure 5-2.)

Lamp: GTE Sylvania 120 MB 6W, Ohmeda Medical Stock No. 0690-2100-315.

1. Disconnect the Warmer power cord and allow the unit to cool for 10 minutes.

2. Use a Phillips head screwdriver to remove the lens mounting screw located in the center of the alarm light.

3. Remove the bulb by pushing in and turning it counterclockwise.

4. Install the new bulb by pushing in and turning it clockwise.

   Note: Replace both bulbs for maximum lamp life.

5. Place the lens cover in position and secure it with the mounting screw.

6. Plug the power cord in and check for proper operation.

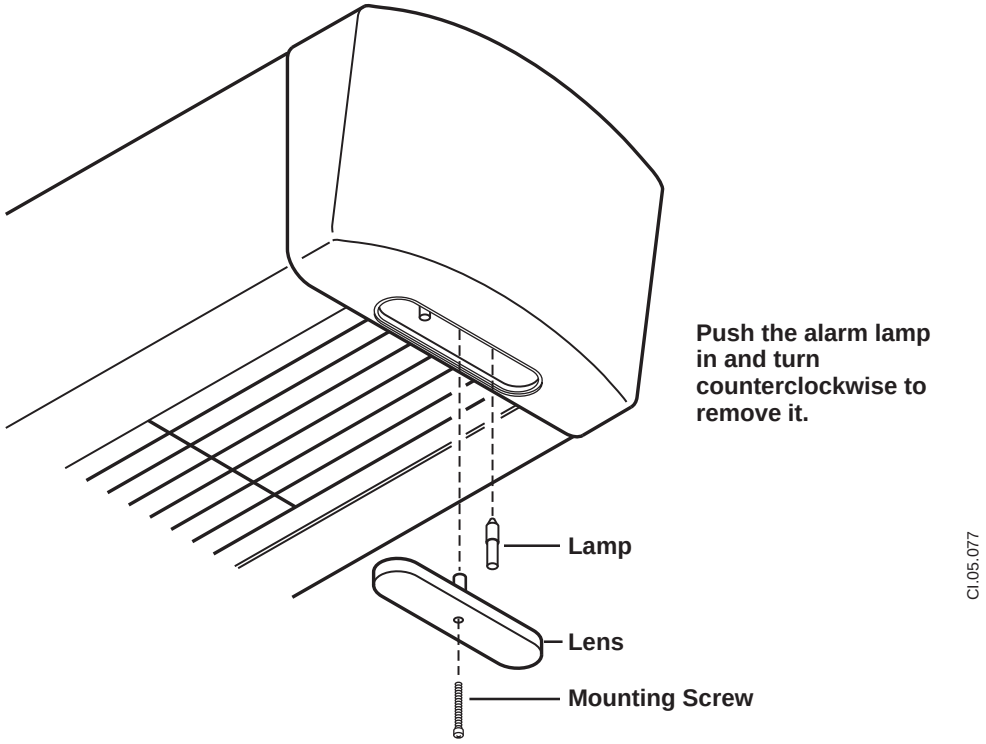

Figure 5-2
Alarm Lamp replacement

B. Observation Lamp Replacement

(See Figures 5-8, 5-9 and 5-10.)

Lamp: GE EXZ (Q50 MR16/NFL), Ohmeda Medical Stock No. 0208-0516-300 or GE EXN (Q50 MR16/FL).

1. Disconnect the power cord for the Warmer and allow the unit to cool for 10 minutes.

2. Rotate the heater assembly to the side position (shown in Figure 5-3).

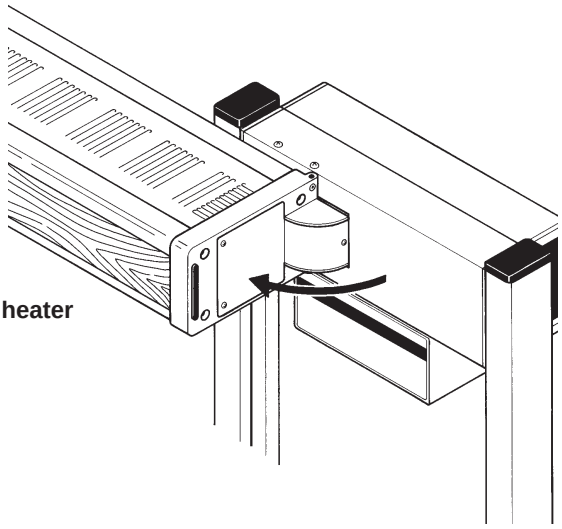

Figure 5-3
Heater rotation to the side position

3. Refer to Figure 5-4. Use a Phillips head screwdriver to remove the back panel of the heater assembly.

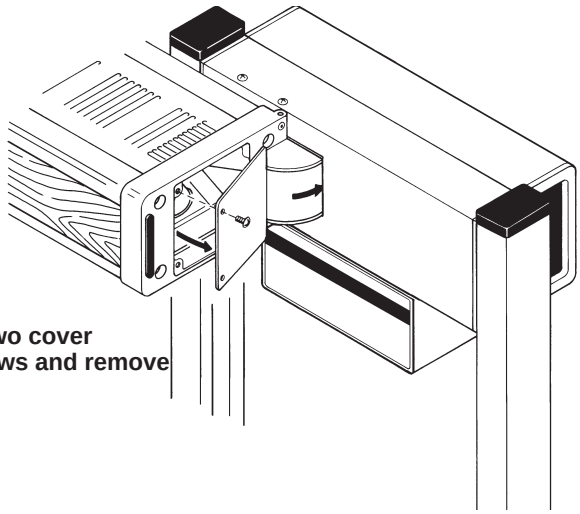

Figure 5-4
Observation Lamp cover removal

4. Refer to Figure 5-5. While holding the lamp with one hand, use your other hand to pull the lever next to the lamp toward you. Remove the lamp.
5. Place the new lamp in position and push it into the lamp socket.
6. Replace the cover and the mounting screws.
7. Rotate the heater assembly back to its normal operating position.
8. Plug the power cord in and switch the observation lamp on. Check for proper operation.

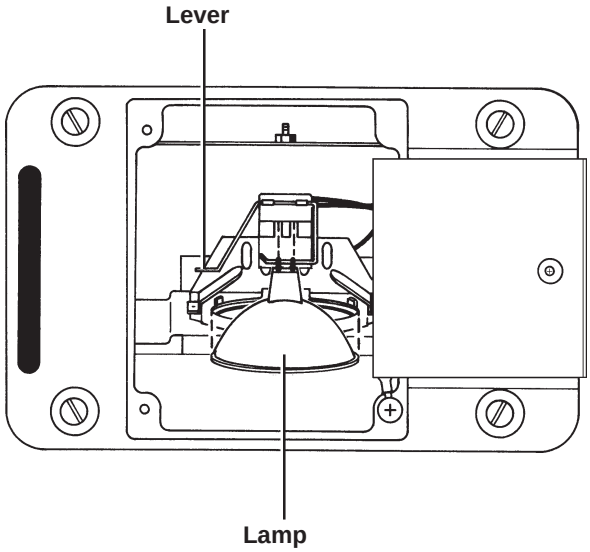

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Figure 5-5
Observation Lamp replacement

5.4 Yoke Manifold Assembly

WARNING: ⚠ **Never oil or grease oxygen equipment. Oils and grease oxidize readily, and in the presence of oxygen, will burn violently. Vac Kote is the oxygen service lubricant recommended (Stock No. 6700-0092-200) if the use of a lubricant is specified.**

Periodically (at least once a year) lubricate the Tee handle screw threads with a small amount of oxygen service lubricant. This prolongs their life and makes sealing of the yoke gaskets easier.

(See Figure 3-9)

Periodically (at least once a year) replace the yoke check valve strainer nipples before they become clogged with lint or dust.

CAUTION: ⚠ Open cylinder valves S-L-O-W-L-Y to avoid damaging regulators.

When installing fresh cylinders, briefly open then immediately close the cylinder valve to blow any foreign matter out of the valve outlet. Do not use excessive force on the shut off valve. Remove the old gasket and use a clean, new gasket (Stock No. 6600-0152-400) in its place.

5.5 Battery Test

The maintenance-free battery should be tested regularly and replaced if necessary. Refer to Section 2.3 for testing the battery. The battery has a two year replacement schedule. Refer to the Service Manual for battery replacement.

5.6 Maintenance Schedule

The unit should be maintained in accordance with the procedures detailed in the Service Manual. Service maintenance must be performed by a technically competent individual as described in the Repair Policy.

A. Operator Maintenance

This schedule lists the minimum frequencies. Always follow hospital and local regulations for required frequencies.

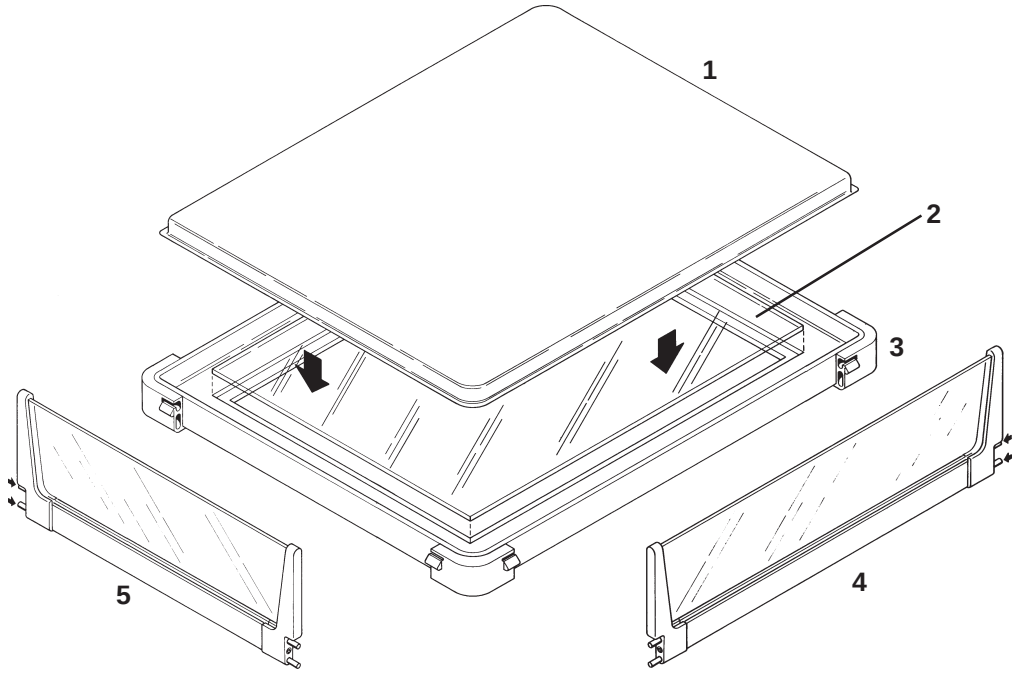
Weekly or after each patient	Clean the Warmer.
After each gas cylinder replacment	Discard old gasket. Use a clean, new gasket.
Annually	Lubricate the Tee handle screw threads (On models equipped with gas package).

B. Service Maintenance

This schedule lists the minimum frequencies. Always follow hospital and local regulations for required frequencies.

Per hospital protocol	Perform the electrical safety and checkout procedure from the Service Manual.
Annually	Check the calibration and perform the service checkout procedure as detailed in the Service Manual.
Annually	Inspect the yoke strainer nipples and replace if necessary (On models equipped with gas package).
Every two years	Replace the battery.
Every five years	Replace the regulator(s) (On models equipped with gas package).

Figure 6-1
Illustrated parts



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Stock Number		Description
0305-5060-300	1.	Small mattress 46.2 x 64 cm
0305-5061-300		Large mattress 59 x 74.2 cm
0217-5221-300	2.	Mattress Support Panel - all models except 3500
6600-0228-800	3.	Corner Block Kit, black - all models except 3500 (4 Blocks)
6600-0230-800		Corner Block Kit, black - model 3500 (4 Blocks)
6600-0185-700		Corner Block Kit, black - model 3500 (Single Block)
6600-0228-801		Corner Block Kit, gray - all models except 3500 (4 Blocks)
6600-0230-801		Corner Block Kit, gray - model 3500 (4 Blocks)
6600-0185-701		Corner Block Kit, gray - model 3500 (1 Block)
6600-0005-800	4.	Left and Right Side Panel, black - model 2001 & series 3000
6600-0009-700		Left and Right Side Panel, black - series 4000
6600-0176-701		Left and Right Side Panel, black - model 3500*
6600-0004-853		Left and Right Side Panel, gray - model 2001 & series 3000
6600-0004-854		Left and Right Side Panel, gray - series 4000
6600-0176-703		Left and Right Side Panel, gray - model 3500*
6600-0004-800	5.	Front and Rear Side Panel, black - model 2001 & series 3000
6600-0008-700		Front and Rear Side Panel, black - series 4000
6600-0176-700		Front and Rear Side Panel, black - model 3500*
6600-0004-855		Front and Rear Side Panel, gray - model 2001 & series 3000
6600-0004-856		Front and Rear Side Panel, gray - series 4000
6600-0176-702		Front and Rear Side Panel, gray - model 3500*

*May be installed as optional, taller side panels on model 2001 and series 3000.

Additional items not shown in Figure 6-1:

6600-0004-801	Tubing organizer panel, black-model 2001 & series 3000 (Figure 3-6)
6600-0004-807	Tubing organizer panel, gray-model 2001 & series 3000 (Figure 3-6)
6600-0004-802	Tubing organizer panel, black-model series 4000 (Figure 3-6)
6600-0004-808	Tubing organizer panel, gray-model series 4000 (Figure 3-6)
6600-0167-716	HFOV Panel, Model 2001, 3300 & 3400 gray
6600-0167-718	HFOV Panel, Model 4300 and 4400 gray
0217-5363-800	Adapter Plate (Figure 6-2)
0217-5372-800	Air Flowmeter with DISS Fittings (50 psi) (Figure 6-3)
6600-0006-800	Air/Oxygen Yoke and Regulator (all model - U.S.)
0217-5377-800	Airway Manometer Assembly (Figure 6-5)
0690-2100-315	Alarm Lamp
6600-0031-900	Bird Dovetail Adapter (includes adapter plate show in Figure 6-2)
6600-0274-800	Chest Drainage Hanger Kit

0212-0300-600	Collection Bottle, 1/4 Gallon
0221-6267-300	Collection Bottle Bracket
0221-6285-300	Collection Bottle Cap with Float Assembly
6702-1224-905	DISS Continuous 3 Mode Suction Regulator with Overflow Trap
6600-0006-850	Drawer Kit, Stationary, beige, 3 drawer (Model 2001 & Series 3000) (Figure 6-11)
6600-0006-851	Drawer Kit, Stationary, beige, 3 drawer (Model 5000 & Series 4000) (Figure 6-11)
6600-0006-852	Drawer Kit, Rotating, beige, 3 drawer (Model 2001 & Series 3000)
6600-0053-850	Drawer Kit, Stationary, gray, 3 drawer (series 3000)
6000-0053-851	Drawer Kit, Stationary, gray, 3 drawer (series 4000)
6000-0053-855	Drawer Kit, Stationary, gray, 2 drawer (series 3000)
6000-0053-856	Drawer Kit, Stationary, gray, 2 drawer (series 4000)
6600-0006-853	Drawer Kit, Rotating, beige (Series 4000)
6600-0053-852	Drawer Kit, Rotating, gray, 3 drawer (Series 3000)
6600-0053-853	Drawer Kit, Rotating, gray, 3 drawer (Series 4000)
6600-0053-857	Drawer Kit, Rotating, gray, 2 drawer (Series 3000)
6600-0053-858	Drawer Kit, Rotating, gray, 2 drawer (Series 4000)
6600-0538-800	ECMO Adapter Kit - model 4400 & 5000
0203-1980-300	Heat Reflective Probe Patch (Box of 50)
0217-5365-800	Instrument Shelf, beige 30 x 30 cm (Figure 6-12)
6600-0513-801	Instrument Shelf, gray 30 x 30 cm (Figure 6-12)
0217-5378-800	IV Pole, 30 cm (Figure 6-4)
0217-5366-800	Monitor Shelf, beige - series 3000 & model 2001, 30 x 66 cm (Figure 6-13)†
6600-0240-801	Monitor Shelf, gray - series 3000 & model 2001, 30 x 66 cm (Figure 6-13)
6600-0240-800	Monitor Shelf, beige - series 4000, 30 x 77 cm (Figure 6-13)
6600-0010-801	Monitor Shelf, gray, - series 4000, 30 x 77 cm (Figure 6-13)
6600-0497-800	Under bed shelf, beige - Model 440
6600-0497-801	Under bed shelf, gray Model 440
0208-0516-300	Observation Lamp
6712-1260-921	Oxygen Flowmeter with DISS Fittings (50 psi)
0304-5260-800	Oxygen Yoke and Regulator Assembly (Figure 6-6 - all models - U.S.)
6600-0055-900	Spot Phototherapy Lamp (wall mounting)
6600-0051-800	Spot Phototherapy Lamp Rail Mounting Kit
0208-0523-300	Power Cord, 120 Volt
0217-5290-870	Routing Clip, black (Package of 6 - Figure 6-7)
6600-0055-851	Routing Clip, gray (Package of 6)
6600-0195-000	Service Manual, all models
0208-0697-700	Skin Temperature Probe Model LA003
6600-0208-700	Skin Temperature Probe Model LA005, Disposable (10/pkg)
6600-0196-700	Skin Temperature Probe Model LA005, Disposable (50/pkg)
6600-0001-850	ThermaLink Upgrade Kit (available for IWS units w/5.0 software or higher)
6600-0004-801	Tubing organizer side panel (rear) - model 2001 & series 3000 † (see Fig. 3-6)
6600-0004-802	Tubing organizer side panel (rear) - model 5000 & series 4000 (see Fig. 3-6)
6700-0136-900	Du-O-Vac
6600-0018-800	Du-O-Vac Mounting Kit
0217-5374-800	Utility Post, 3.5 Inch Stub x 1 in. diameter (Figure 6-8)
0217-5369-800	Vacuum Manifold with DISS Fittings (Figure 6-9)
0217-5367-800	Vacuum Slide Bracket (Figure 6-10)
0217-5357-800	Ventilator Mounting Post Accessory

For more information about these and other accessories see the Infant Care Equipment Accessories Catalog (Form No F058)

†Not compatible with model 3500.

Figure 6-2
Adapter Plate
(0217-5363-800)

Figure 6-3
Air Flowmeter (50 psi) with DISS Fittings (0217-5372-800)

Figure 6-4
IV Pole, 30 cm
(0217-5378-800)

Figure 6-5
Airway Manometer Assembly
(0217-5377-800)

Figure 6-6
Oxygen Yoke and Regulator Assembly (see list on 6-1 and 6-2)

Figure 6-7
Routing Clip
(Package of 6)
(0217-5290-870)

Figure 6-8
Utility Post,
3.5 Inch stub
(0217-5374-800)

Figure 6-9
Vacuum Manifold
with DISS Fittings
(0217-5369-800)

Figure 6-10
Vacuum Slide Bracket
(0217-5367-800)

Figure 6-11
Drawer Module
Stationary
(See list on 6-1 and 6-2)

Figure 6-12
Instrument Shelf
(See list on 6-1 and 6-2)

Figure 6-13
Monitor Shelf
(See list on 6-1 and 6-2)

Appendix

In this section

Temperature Conversion Chart A-1

Servo Mode Algorithm A-2

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Installing wall mounted units A-13

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All specifications are subject to change without notice.

Temperature Conversion Chart

C	F	C	F	C	F
20.0	68.0	34.5	94.1	37.6	99.7
21.5	70.7	35.0	95.0	37.7	99.9
22.0	71.6	35.1	95.2	37.8	100.0
22.5	72.5	35.2	95.4	37.9	100.2
23.0	73.4	35.3	95.4	38.0	100.4
23.5	74.3	35.4	95.7	38.1	100.6
24.0	75.2	35.5	95.9	38.2	100.8
24.5	76.1	35.6	96.1	38.3	100.9
25.0	77.0	35.7	96.3	38.4	101.1
25.5	77.9	35.8	96.4	38.5	101.3
26.0	78.8	35.9	96.6	38.6	101.5
26.5	79.7	36.0	96.8	38.7	101.7
27.0	80.6	36.1	97.0	38.8	101.8
27.5	81.5	36.2	97.2	38.9	102.0
28.0	82.4	36.3	97.3	39.0	102.2
28.5	83.3	36.4	97.5	39.5	103.1
29.0	84.2	36.5	97.7	40.0	104.0
29.5	85.1	36.6	97.9	40.5	104.9
30.0	86.0	36.7	98.1	41.0	105.8
30.5	86.9	36.8	98.2	41.5	106.7
31.0	87.8	36.9	98.4	42.0	107.6
31.5	88.7	37.0	98.6	42.5	108.5
32.0	89.6	37.1	98.8	43.0	109.4
32.5	90.5	37.2	99.0	43.5	110.3
33.0	91.4	37.3	99.1	44.0	111.2
33.5	92.3	37.4	99.3	44.5	112.1
34.0	93.2	37.5	99.5	45.0	113.0

Appendix

Servo Mode Algorithm

The servo mode uses the difference between the servo mode control temperature and the patient skin temperature to determine the percent of heater power required:

Control Temp - Patient Temp			Percent Heater Power
0.45°C		100	
0.44	to	0.30°C	95
0.29	to	0.25°C	90
0.24	to	0.20°C	85
0.19	to	0.15°C	75
0.14	to	0.10°C	65
0.09	to	0.05°C	45
0.04	to	–0.04°C	25
–0.05	to	–0.09°C	20
–0.10	to	–0.14°C	15
–0.15	to	–0.19°C	10
–0.20	to	–0.24°C	5
–0.25°C			0

Infant Warmer System Specifications

Electrical

Power Requirements:

120 V, 50/60 Hz Model: 115 V ~ ± 10%, 6.6 Amps.

220 V, 50/60 Hz Model: 220 V ~ ± 10%, 3.7 Amps.

230 V, 50/60 Hz Model: 230 V ~ + 10%, –6%, 3.4 Amps.

240 V, 50/60 Hz Model: 240 V ~ ± 10%, 3.3 Amps.

100 V, 50/60 Hz Model: 95 V ~ ± 10%, 8.2 Amps.

All models designed to conform to BSI 5724, IEC 601-1, UL544, CSA 22.2, SEMKO and TUV requirements.

Heater Output:

All models:

540 watts ± 5% at maximum % power setting.

Average Energy at Mattress Level at maximum % power setting:

20 mw/cm^2 for Model 5000 & Series 4000.

34 mw/cm^2 for Model 2001 & Series 3000.

Recommended Bed Level:

69±5 cm from bottom of heater module.

WARNING: ⚠ **Bed-to-heater spacing less than the recommended 64 cm will result in incorrect operation and may adversly affect the patient's condition**.

Appendix

For your convenience, on Model 3000, 3100 and 3500 the proper bed-to-heater spacing is indicated by a label located on the right aluminum upright (as viewed from the front).

Line Voltage Compensation:

Input line voltage is monitored and the heater drive is adjusted to compensate for variations in the line voltage. This ensures a stable heater output despite input line voltage fluctuations.

Circuit Breaker:

Rated Current: 7.5 A.

Trip Point: 9.45 A Minimum.

Type: Manual Resetting.

Model: Airpax Snapak.

Isolation Voltage:

2500 Vrms 60 Hz from the patient probe to the ac phase and neutral lines for one minute.

WARNING: ⚠ **The patient probe is not isolated from earth ground. Any additional equipment used with the Ohmeda Infant Warmer System must comply with UL 544, CSA 22.2, IEC 601, and VDE 750.**

Leakage Current:

With ground wire open or connected and measured at an exposed metal surface, less than 100 microamperes on 100 V and 120 V units (200 microamperes on 220 V, 230 V and 240 V units).

Ground Resistance:

Less than 0.2 ohms as measured from the power cord plug

IEC 601-1 Specifications

Type of protection against electric shock:	Class 1
Degree of protection against electric shock:	Type B
Mode of Operation:	Continuous
Protection against hazards of explosion:	Not Protected
Protection against ingress of liquids:	Not Protected
Degree of electrical connection between equipment and patient:	Equipment with an applied part designed for non-conductive connection to the patient.
Degree of mobility:	Fixed (Model 3050, 3100 and 3150) Mobile (all others)

Controller

Electronics:

Microprocessor-based control system.

Self-test functions are performed at power up and during normal operation.

Power Control Method:

Proportional heat control with zero-voltage switching to minimize radiated and conducted EMI.

Observation Light:

Nominal illuminance output: 100 foot candles at center of mattress.

Estimated lamp life: 3,000 hours.

Temperature Sensing System:

Range: 30 - 42°C
Accuracy: ±0.3°C
Resolution: ±0.1°C
Probe interchangeability: ±0.1°C
Probe Model Number: LA003 or LA005

Elapsed Timer:

60 minute elapsed timer with hold mode and Apgar tones.

Manual Mode Heat Selector Range:

All models:

0 to 540 watts in 20 increments of 5% each.

Servo Mode Control Range:

(all models except the 2001 international)

35.0 to 37.5°C in increments of 0.1°C.

Alarms

Multiple audio tones:

Operator prompt tone.*
Alternating single tone.*
Alternating two tone.*
*For a more detailed description of the audible alarms see section 3.1D

Overhead Alarm Light:

Large alarm light located on the front of the heater assembly for easy visual identification.

Probe Failure Alarm:

The alternating two tone alarm is active only in the servo mode.

Activates when the skin temperature probe:

1. Fails electrically due to an open or short circuit, or
2. Is disconnected from the Warmer.
3. Probe senses temperature outside the 30 - 42°C range.

When this alarm condition exists:

1. The heater is turned off and
2. The patient temperature display flashes “HH.H” or “LL.L”.

Patient Temp. Alarm:

This single tone alarm activates in the servo mode when the difference between the patient temperature and the control temperature is greater than 1°C (can be adjusted to 0.5°C by qualified service person).

Alarm cancels when the patient temperature returns to within 0.8°C of the control temperature.

System Failure Alarm:

This alternating, two tone alarm cannot be silenced.

Alarm activates and turns the heater off if any of the following occurs:

1. The analog-to-digital converter calibration drifts by more than 0.3°C.

2. The heater solid state relay fails.

3. The microprocessor fails or -

4. The self check functions fail on power-up.

Note: Excessive EMI in the hospital environment can trigger the system failure alarm. Note the error code, if any, and switch off the unit. Wait ten seconds and switch the power back on. If the system fail alarm recurs, remove the warmer from use.

Check Patient Alarm:

Manual Mode: Single tone alarm activates if the heater has been energized at greater than 25% heat for 12 continuous minutes.

Servo mode: Alarm activates when the heater has been at full power for 12 continuous minutes.

Alternating two tone alarm activates after 3 minutes if the Check Patient Alarm is not silenced.

Power Failure Alarm:

Single tone alarm activates if the line power is interrupted. The power failure LED on the display panel lights.

A rechargeable maintenance-free nickel cadmium battery powers the audio alarm and the microprocessor for up to 10 minutes with a fully charged battery. If power is re-stored within 10 minutes, the mode of operation and the set point are recalled.

Heat Off Alarm:

The Light-Emitting-Diode (LED) indicator activates when the heater is in the side position.

The audio alarm activates after the heater has been in the side position for 5 minutes.

Environmental Specifications

Operating Temperature Range: 10 to 40°C.

Transport and Storage Temperature Range: -25 to 60°C.

Transport, Operating and Storage Pressure Range: 500 to 1060 hPa

Operating and Storage Relative Humidity Range: 0 to 95%.

Electromagnetic Compatibility (EMC) Specifications

All models meet the requirements as specified in IEC 601-1-2
Collateral Standard Electromagnetic Compatibility - Requirements and Tests.

Mechanical (Without Accessories)

Overall Dimensions:

Model number	Height - cm	Depth - cm	Width - cm	Minimum Wt** kg
2001	185	104	66	96
3000	185	114	66	70
3050	28*	84	46	18
3100	152*	84	55	34
3150	152*	91	55	50
3300	185	104	66	96
3400	183-203	114	77	91
3500†	185	103	62	102
4000	185	132	77	80
4300	185	114	77	101
4400	183-203	114	77	91

* Height of equipment itself, height from floor will vary on application.

† Includes bassinet.

**Drawers not included; for stationary drawers add 25 kg, for rotating drawers add 32 kg.

Mattress dimensions:

Model number	Width - cm	Length - cm
2001	46	64
3000	—	—
3050	—	—
3100	—	—
3150	46	64
3300	46	64
3400	47	64
3500	46	64
4000	—	—
4300	59	74
4400	59	74

Casters

Model number	
2001	13 cm dia., 2 locking, 2 non locking
3000	13 cm dia., 2 locking, 2 non locking
3050	—
3100	—
3150	—
3300	13 cm dia., 2 locking, 2 non locking
3400	13 cm dia., 2 locking, 2 non locking
3500	Warmer- 5 cm dia., 2 locking, 4 non locking Bassinet- 13 cm dia., 2 locking, 2 non locking
4000	13 cm dia, 2 locking, 2 non locking
4300	13 cm dia., 2 locking, 2 non locking
4400	13 cm dia., 2 locking, 2 non locking

Drawers†

Model number	
2001	3 drawers- 39 x 39 x 10 cm or 1 drawer 39 x 39 x 10 cm, 1 drawer 39 x 39 x 23 cm
3000	—
3050	—
3100	—
3150	3 drawers- 38 x 39 x 10 cm or 1 drawer 38 x 39 x 10 cm, 1 drawer 38 x 39 x 23 cm
3300	3 drawers- 38 x 39 x 10 cm or 1 drawer 38 x 39 x 10 cm, 1 drawer 38 x 39 x 23 cm
3400	3 drawers- 38 x 39 x 10 cm or 1 drawer 38 x 39 x 10 cm, 1 drawer 38 x 39 x 23 cm
3500	1 drawer- 30 x 39 x 8 cm 2 drawers - 39 x 39 x 15 cm
4000	—
4300	3 drawers- 38 x 39 x 10 cm or 1 drawer 38 x 39 x 10 cm, 1 drawer 38 x 39 x 23 cm
4400	3 drawers- 38 x 39 x 10 cm or 1 drawer 38 x 39 x 10 cm, 1 drawer 38 x 39 x 23 cm

† Dimensions for both the stationary and rotating drawer packages are the same.

Bed duty cycle (Elevating models only)

1 minute on

10 minutes off

Bed tilt positions

3500	± 8° in increments of 4°
All other models	± 10° continuously adjustable

3500 Bassinet

Material:

Select oak and oak veneer. Optional custom stain colors are available.

Storage:

Three Drawer storage opening from either side or the front of the unit, depending on the model ordered.

Accessories

Accessories include the following (see also Section 6/Illustrated Parts):

Air/Oxygen Yoke and Regulator:

Pin indexed oxygen yokes accommodate two U.S. E size oxygen cylinders.

Additional yoke accommodates one U.S. E-size cylinder of compressed air.

Air and oxygen pipeline fittings.

Regulators: 358 ± 14 kPa.

Cylinder pressure gauges: 0 to 20700 kPa.

Oxygen Yoke and Regulator:

Pin indexed yokes accommodate two U.S. E size oxygen cylinders.

Oxygen pipeline fittings.

Regulator: 358 ± 14 kPa.

Cylinder pressure gauge: 0 to 20700 kPa.

Rail Mounted Accessories:

WARNING:

⚠ **Overloading the shelves can affect the stability of the unit. Limit the load to 9 kg per instrument shelf, mounted to a single upright, and 23 kg per monitor shelf, mounted between the uprights.**

⚠ **Limit the load of accessories to 23 kg per side on the Warmer to ensure stability. Always try to evenly distribute the weight of accessories on both sides of the unit for amore balanced load. Accessories should not be mounted more than 56 inches (142 cm) above the floor. For models 3000 and 3500, limit the load of accessories to 9 kg maximum per side mounted no more than 112 cm above the floor.**

⚠ **Due to the increased height of units with the ECMO option installed, a tipping hazard may exist. Limit the total accessory load to 23 kg, no more than 11 kg per side.**

System 2001/3000/3300 Monitor Shelf:

Dimensions 30 x 58 cm
Load limit: 22 kg

System 4000 Monitor Shelf:

Dimensions 30 x 77 cm
Load Limit: 22 kg

Instrument shelf:

Dimensions 30 x 30 cm
Load limit: 9kg

Oxygen flowmeter with DISS fittings: 0 to 15 LPM

Air flowmeter with DISS fittings: 0 to 15 LPM

Airway Manometer: -20 to +100 centimeters of water

IV pole - 30 and 60 cm

Gas manifold with 1/8 inch NPT fitting

9 cm utility post

Drawer Storage Accessory:

(for all models with beds - stationary or rotating)

Standard drawer 38 x 39 x 10 cm

14,820 cm^3 per drawer

Deep Drawer 38 x 39 x 23 cm

34,086 cm^3 per drawer

WARNING: ⚠ **Overloading the drawers can affect the stability of the unit. Limit the load to 4.6 kg per drawer, 9 kg per deep drawer.**

Utility Post, 22 Inch (0217-5376-800)

2.5 cm diameter x 56 cm long post

Provides a rigid vertical mounting space for stack mounting of equipment. The post should be mounted as low as possible to ensure the stability of the Warmer.

Vacuum Manifold with DISS Adapters

Two DISS vacuum adapters are mounted on a standard manifold block. The locations of these adapters can be changed to any of the other tapped holes in the block to meet special requirements.

Vacuum Bottle Slide Bracket

The standard vacuum bottle slide can be mounted on either upright.

Ventilator Mounting Post

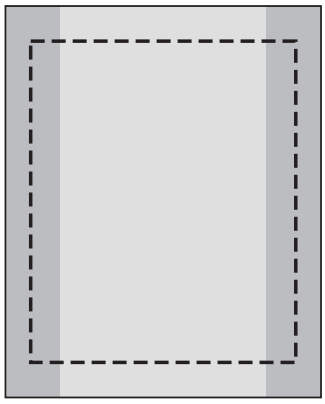
The vertical mounting post is a non-locking 1 inch diameter shaft which pivots in an 22 cm radius from the upright. One end of the shaft extends 36 cm while the other extends 16 cm. Ventilators, humidifiers, blenders or other user hardware can be pivoted into optional position.

Radiant Energy Distribution

This table lists typical average radiant energy distribution across the patient bed surface for informational purposes only.

	mW/cm²	
	Zone Average	Mattress Average
Model 2001 and Series 3000 units†	34	34
Series 4000 units	22	20

Typical Radiant Energy Level on the Mattress Surface



†The radiant energy level values of older models will be less than those shown; units prior to those with HCA serial numbers had a 440 Watt heater.

CI.14.001

Infant Zone - 15" (38 cm) wide

Mattress - Model 2001/Series 3000

Mattress - Series 4000

Thermalink Option Specifications

Hewlett Packard CMS Monitor (Vuelink)

The warmer may be connected to a Hewlett Packard CMS Monitor via a Vuelink open interface module (HP P/N M1032A Option A05) and HP Ohmeda option cable. The parameters passed to the HP monitor are: patient temperature, set temperature, and heater power level; these three parameters are displayed by default on power-up. The user can select to display any or all of these parameters. All alarms are also passed to the monitor. See the Hewlett Packard CMS and Vuelink manuals for features and functions of the HP system.

SpaceLabs Monitor (Flexport)

The warmer may be connected to a SpaceLabs Bedside Monitor via a Flexport (SL P/N 90444A). The parameters passed to the SpaceLabs Bedside Monitor are: patient temperature, set temperature, and heater power level. All alarms are also passed to the monitor. See the SpaceLabs Monitor and Flexport manuals for features and functions of the SL system

Serial data

RS-232 Connector

WARNING: ⚠ **The computer or RS-232 monitor's user program must continuously check the data link. The program should constantly verify connection to the warmer controller and check for updated data.**

Note: In the event of power failure, all serial communication will cease until power is restored.

The Nurse Call and the serial data output share the same female, nine pin, d-type connector (DB9F).

Pin 2: Receive Data (warmer input)
Pin 3: Transmit Data (warmer output)
Pin 5: Gnd (Signal Ground)

Cable requirements

The user interface cable must have capacitance less than 1500 pF. It should be a shielded cable such as a Belden 9611 with AMP shielding kit 748046-1 and ferrule 747579-8.

Data transmission

The warmer continuously sends data from the time that it is first powered up. Note that the warmer serial data transmission can be controlled through the serial port. Data output stops when the warmer receives a <cntrl>S (XOFF) and resumes when it receives a <cntrl>Q (XON).

Data format

1 start bit, 7 data bits (ASCII), 1 parity bit (odd), 1 stop bit, 1200 baud, full duplex.

Serial data has the format: start text character, “<stx>”; IWS header, “IW”; software version; data string; checksum characters; carriage return, “<cr>”; line feed, “<lf>”; end of text character, “<etx>”. Data elements are separated by spaces, “_”. Each String contains 53 characters:

Sample data:

<stx>IW0500_35.52_XX.XX_P_00085_36.50_00001000_E014_11<cr><lf><etx>

Data for discussion (use the following table)**:**

<stx>IWxxxx_pt.pt_xx.xx_m_ppppp_pc.pc_alrmleds_code_ck<cr><lf><etx>

<stx>

Start of text character (ASCII 2) or CTRL B; indicates a string of data will follow.

IWxxxx

IW means the data is from the Infant Warmer; xxxx is the software version in the unit, e.g. 0500 for version 5.00.

pt.pt

This is the patient temperature in degrees centigrade. The patient temperature will always be sent even if it is outside the normal display range. Temperatures less than or equal to 1.00°C indicate an open or a disconnected probe. Any temperature greater than or equal to 50°C is a shorted probe.

xx.xx

Reserved for future use.

m

This is the mode of operation. P means servo control mode. A means manual control mode. Always check the mode of operation before evaluating the patient control temperatures.

ppppp

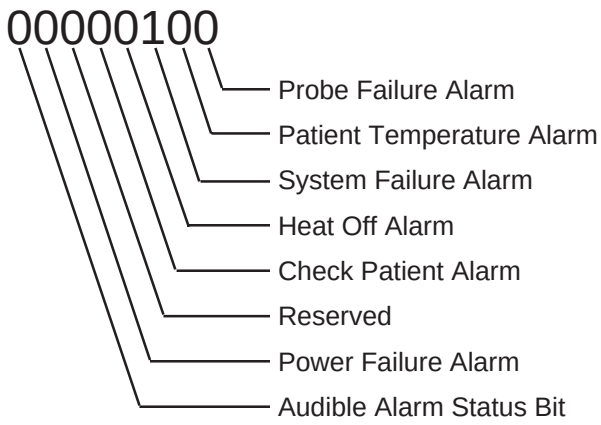
Percentage of nominal power defined as % of heater wattage.

pc.pc

In the servo control mode, this is the patient control temperature set with the ▲ and ▼ switches in degrees centigrade.

alrmleds

This series of bits represents the alarm LEDs. If an LED is illuminated, the corresponding bit is set to 1 (alarm active). If there is no alarm, the bit is set to 0. Checking the bits from right to left is the same as checking the alarm LEDs from top to bottom:



code

This is the error code that appears in the control temperature display during a system failure alarm (e.g. E014). If the system is operating normally (no system failure), zeroes replace the error code (e.g. E000).

ck

This is the two byte ASCII representation of the byte that when added to the sum of all the ASCII data bytes in the string equals zero. Note that all over flows are dropped and the sum of the data bytes DOES NOT INCLUDE the <stx>, checksum, <etx>, <cr>, or <lf> characters or the parity bit of each byte.

<cr>

Carriage return character.

<lf>

Line feed character.

<etx>

End of transmission character (ASCII 3) or CTRL B.

Nurse Call specifications

Contact ratings

Maximum resistive load: 4 VA
Maximum DC switching voltage: 100 Vdc
Maximum switching current: 0.25 A
Maximum carrying current: 0.50 A.

Connector

The Nurse Call contacts and the serial data output share the same female, nine pin, d-type connector (DB9F).

Pin 6:	Closed contact under normal conditions, i.e. power on, no alarm (recommended configuration)
Pin 1:	Common contact
Pin 9:	Open contact under normal conditions, power on, no alarm

These contacts are not powered. They only provide closure.

Appendix

Table A-1
Nurse Call signals

Warmer Status	Nurse Call Signal Pins 1&6	Pins 1&9
Normal	Closed	Open
Alarm	Open	Closed
Power switch off or power fails	Open	Closed
Nurse Call cable disconnected	Open	Open

WARNING: ⚠ **If you use the normally open Nurse call connection, a disconnected Nurse Call cable DOES NOT trigger a remote alarm.**

Note: Any interruption of warmer power (deliberately switching off the warmer , accidentally unplugging the power cord, etc.) triggers a Nurse Call alarm.

Installing wall mounted units

WARNING: ⚠ **Carefully follow these installation instructions. Failure to do so may result in serious injury to the operator or patient.**

The model 3050, 3100, and 3150 wall mounted warmers were designed for locations where permanent, infant radiant warmers attached directly to the room wall are desired. To adequately install these models, pre-planning at the architectural design phase is required.

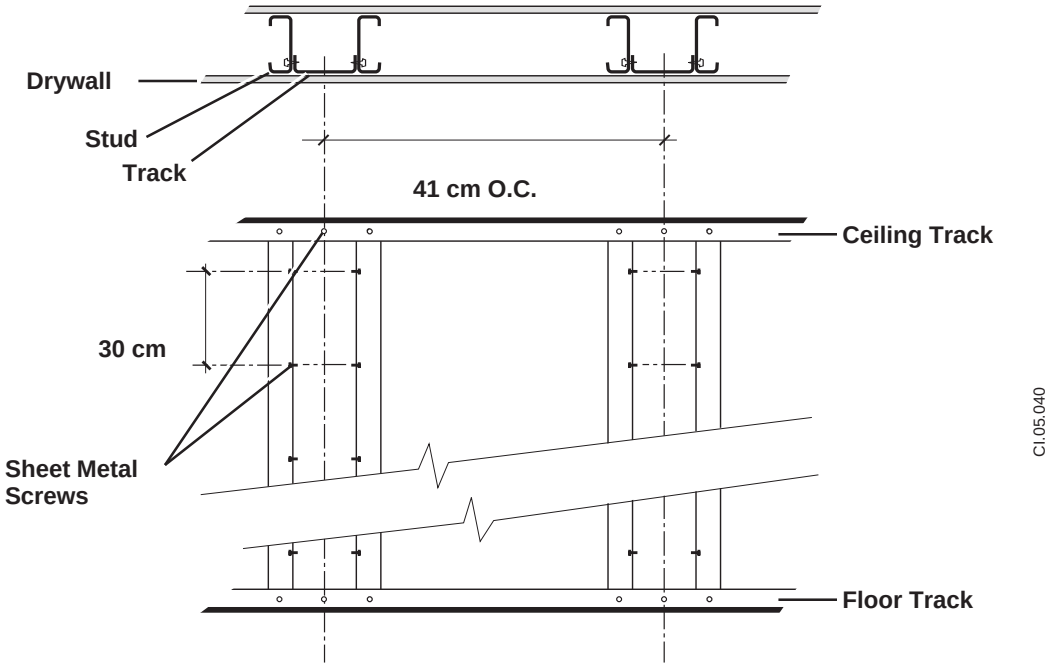


Figure A-1
Wall cross section

The following instructions describe the manufacturer's recommended method of installation in new construction. Use only the Ohmeda hardware provided to mount the model 3050, 3100, and 3150. The installation should be approved by the appropriate State and Local authorities. Deviation from these recommendations or attempts to retrofit existing construction should only be undertaken by professionals experienced in structural design, who in turn are wholly responsible for the structural integrity of the mounting method they devise.

Pre-Installation Preparation

A pattern of two (2) Extra Duty Drywall Studs (STE-20 gauge) and one (1) 10 x 4 cm Drywall Track (16 gauge) at 41 cm centers form the basic "rough in" for installing each IWS 3050 unit. See Figure A-1. Verify that the studs and track are securely fastened together at 30 cm intervals and to both the floor and ceiling tracks with #10 sheet metal screws.

Warmer Installation

After the wall is finished and the room is completed you are ready to install the warmer.

The distance from the warmer heater head to the bassinet or bed mattress surface is critical for proper operation of the warmer. This distance must be 69 - 5, +23 cm. Before installing the model 3050 or 3100, you must determine the bed surface height that will be used (for your convenience, a bed height label is provided to attach to the wall to aid in visually aligning the bassinet/bed surface).

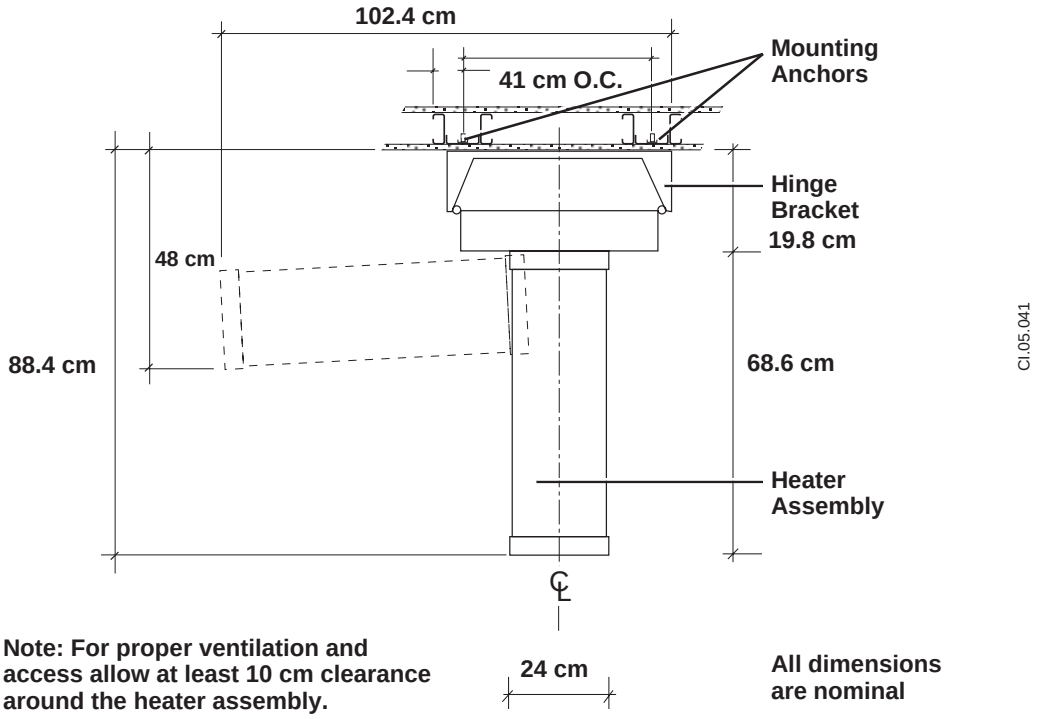


Figure A-2
Warmer mounitng overhead view

1. Locate the stud edges by pre-measurement or by using a stud finder. Find the center of the drywall track 86 mm over from the stud edge. See Figure A-2.

2. Remove the hinge bracket from the warmer by removing the (2) hinge pins (one on either side) that secure the bracket to the warmer heater head.

3. Utilizing a level, position the hinge bracket on the wall. The bottom edge of the heater mounting bracket on the model 3050 and 3100 must be 69 - 5, + 23 cm, inches from the bed surface of the mattress that will be used. See Figure A-3. The model 3150 integral bed warmer already comes with 69±5 cm bed-to-heater spacing.

4. Use the hinge bracket as a template to mark the mounting holes on the wall.

5. Use a 19 mm diameter metal cutting hole saw to drill (4) holes through the dry wall and through metal drywall tracks behind the wall.

WARNING:

⚠ **When installing wall mounted units, to achieve adequate structural strength all four holes must engage tracks. Should the hole saw miss a track do not proceed. Consult the project engineer for further direction.**

⚠ **Enclosing the heater assembly inside a cabinet may prevent proper ventilation and may create a fire hazard. If the heater is enclosed in a cabinet, it must be equipped with a power cut-off device that prevents operation while the cabinet is closed.**

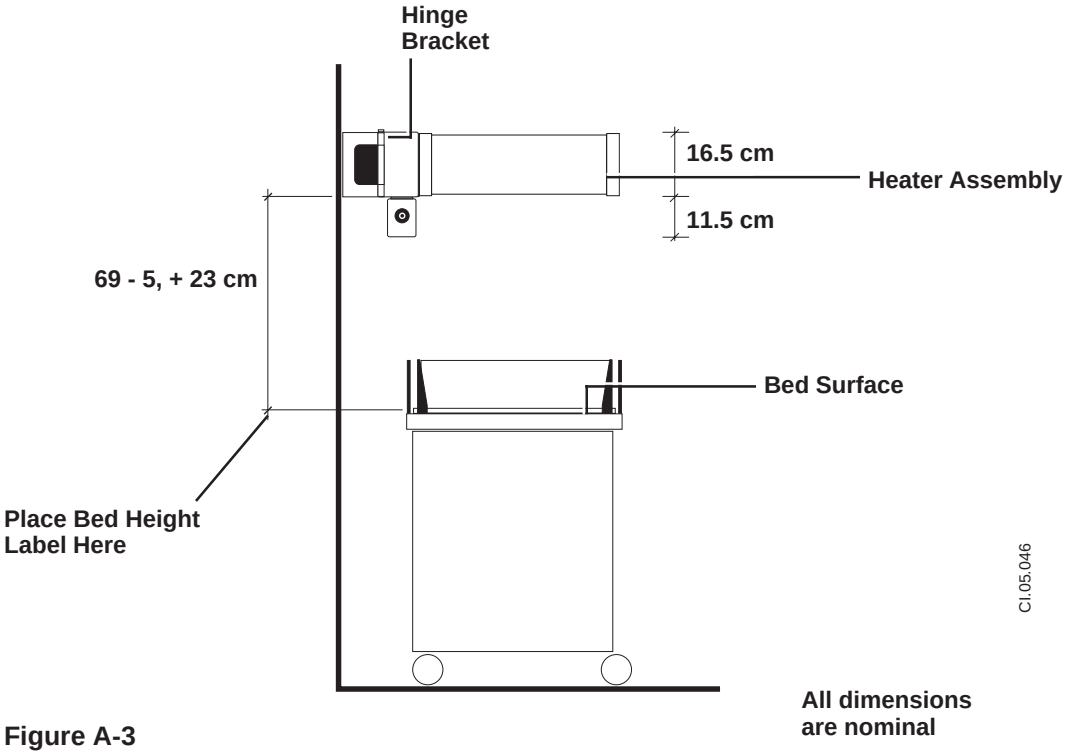


Figure A-3
Heater to bed distance

6. Perform steps A through C to secure the (4) Toggler® anchor bolts in the wall. See Figure A-4.

   A. Hold the metal channel flat alongside the plastic straps and slide the channel through the hole in the wall and track.

   B. With one hand, hold the ring so the metal channel rests flush behind the wall. Slide the plastic cap along straps with the other hand until the flange of the cap is flush with the wall.

   C. Place your thumb between the straps, and push from side to side, snapping off the straps level with the flange of the cap.

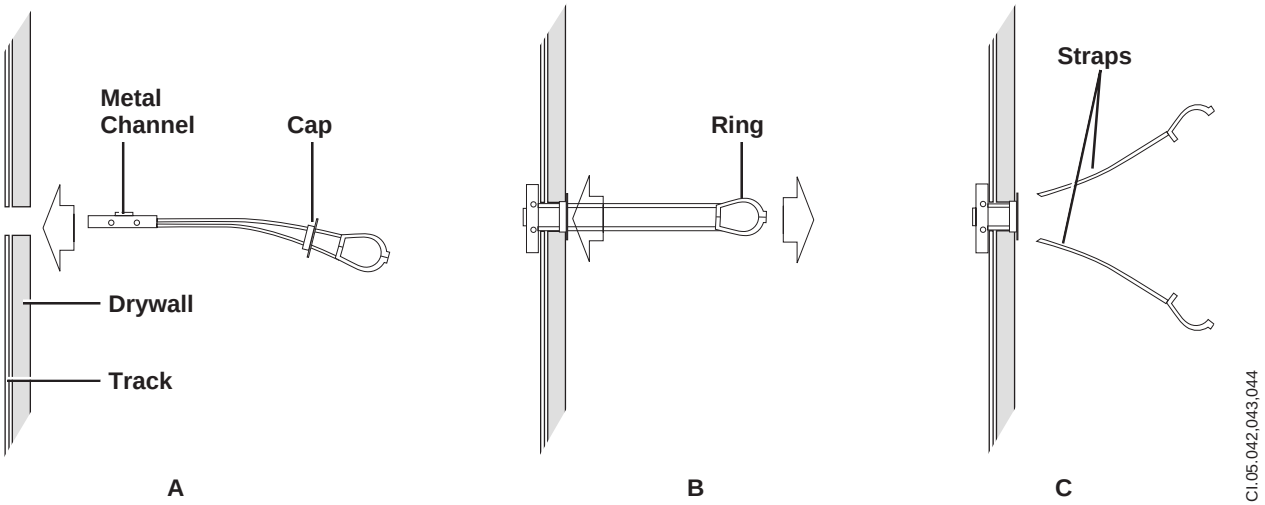


Figure A-4
Toggler® anchor installation

7. Mount the hinge bracket on the wall with the flat washers, split ring washers and screws provided. See Figure A-5. Tighten the screws with approximately 80 lb/in of torque.

   Note: The screws provided (3/8 - 16 UNC, 2 ½" long) are for installations in 1/2 inch or 5/8 inch thick drywall. For applications involving double thicknesses of drywall, longer screws will be required. To properly mount the warmer, at least 2 a screw threads must be engaged.

   Verify that the bracket is level and the distance to the bed surface is 69 - 5, + 23 cm.

WARNING: ⚠ **Keep hands clear of the hinge area when installing a wall mounted heater assembly. A pinch hazard exists.**

⚠ **Heater assembly weighs approximately 14 kg. Proper installation may require two people. Due to the weight of the 3100 and 3150 units, approximately 34 kg, proper installation will require two people.**

8. Mount the warmer by inserting the back of the unit into the hinge bracket mounted on the wall, then securing it by tapping in the (2) hinge pins. Verify that the heater assembly is rigidly secured to the wall, is level, and is parallel to the floor.

9. For the model 3050, attach the "proper bed to heater spacing" label provided with the unit to the wall so that the bottom line of the label is 74 cm from the lower edge of the hinge bracket.

WARNING: ⚠ **Bed-to-heater spacing less than the recommended 64 cm will result in incorrect operation and may adversly affect the patient's condition.**

For the model 3100 or 3150 perform steps 10 through 14. For the model 3050 skip to step 15.

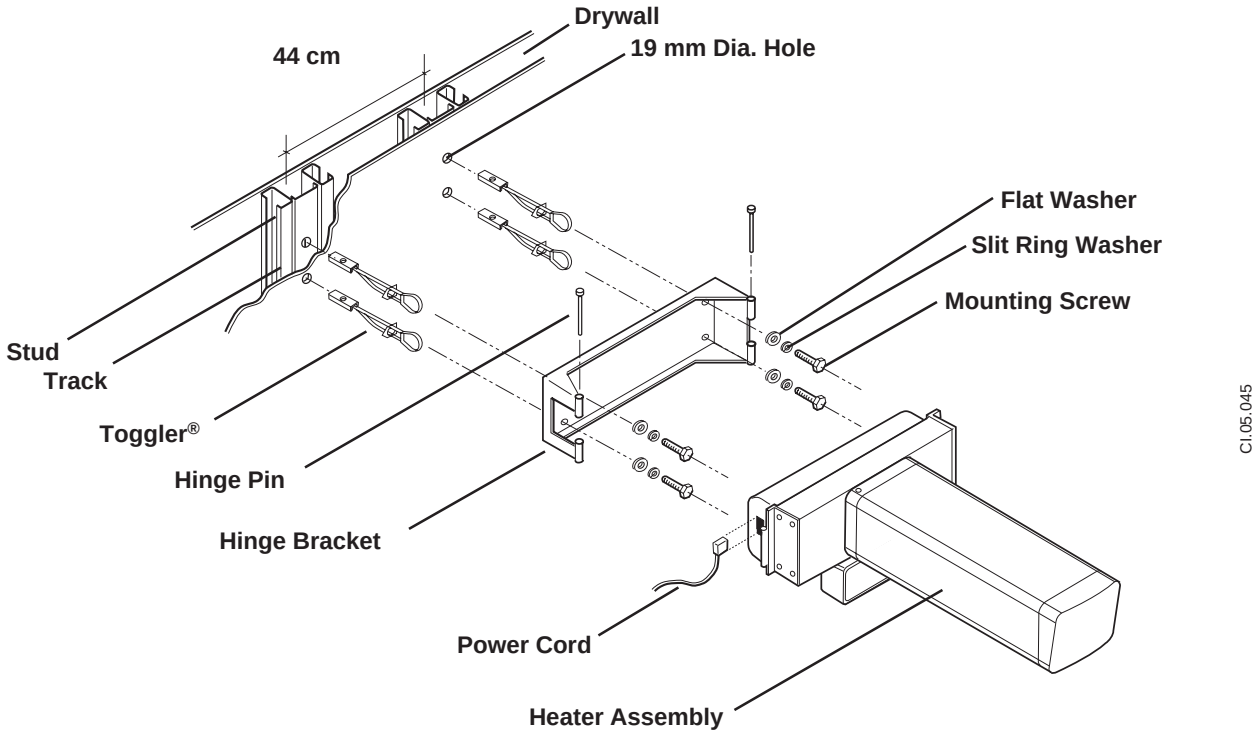


Figure A-5
Mounting the warmer

10. Position the lower mounting bracket on the wall. Use the lower hinge bracket as a template to mark the mounting holes on the wall for the lower bracket.

11. Remove the warmer from the wall by tapping out the (2) hinge pins from the upper bracket.

12. Remove the lower hinge bracket from the warmer by removing the (2) hinge pins (one on either side) that secure the bracket to the warmer.

13. Repeat steps 5, 6 and 7 to properly install the lower mounting bracket.

14. Mount the warmer by inserting the back of the unit into the upper hinge bracket mounted on the wall, then securing by tapping in the (2) hinge pins. Secure the warmer to the lower mounting bracket by tapping in the (2) hinge pins. Verify that the heater assembly and side rails are rigidly secured to the wall and that the heater is level and is parallel to the floor.

15. Refer to the Operation & Maintenance Manual for checkout procedures.

WARNING: ⚠ **Perform Checkout Procedures before placing the unit in operation.**

Notes

Notes

Notes

Warranty

This Product is sold by Ohmeda Medical under the warranties set forth in the following paragraphs. Such warranties are extended only with respect to the purchase of this Product directly from Ohmeda Medical or Ohmeda Medical's Authorized Dealers as new merchandise and are extended to the first Buyer thereof, other than for purpose of resale.

For a period of twelve (12) months from the date of original delivery to Buyer or to Buyer's order, but in no event for a period of more than two years from the date of original delivery by Ohmeda Medical to an Ohmeda Medical Authorized Dealer, this Product, other than its expendable parts, is warranted to be free from functional defects in materials and workmanship and to conform to the description of the Product contained in this operation manual and accompanying labels and/ or inserts, provided that the same is properly operated under conditions of normal use, that regular periodic maintenance and service is performed and that replacements and repairs are made in accordance with the instructions provided. This same warranty is made for a period of thirty (30) days with respect to the expendable parts, including but not limited to the reusable Ohio skin temperature probe. The foregoing warranties shall not apply if the Product has been repaired other than by Ohmeda Medical or in accordance with written instructions provided by Ohmeda Medical, or altered by anyone other than Ohmeda Medical, or if the Product has been subject to abuse, misuse, negligence, or accident.

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OHMEDA Medical

Ohmeda Medical
8880 Gorman Road
Laurel MD 20723
USA
410-888-5200
Fax 410-888-0544

North America

United States

Customer Service and Distribution Center
Datex-Ohmeda, Inc.
Ohmeda Drive
PO Box 7550
Madison, WI 53707-7550
Tel 1 800 345 2700
Fax 1 608 221 4384

Technical Support
Datex-Ohmeda, Inc.
Three Highwood Drive
Tewksbury, MA 01876
Tel 1 800 345 2755

Sales and Service
Datex-Ohmeda, Inc.
Three Highwood Drive
Tewksbury, MA 01876
Tel 1 800 635 6099
Fax 1 978 640 0469

Equipment Service Center
Datex-Ohmeda, Inc.
1315 West Century Drive
Louisville, CO 80027-9560
Tel 1 800 345 2755

Canada

Dynamed Health Care Systems
235 Shields Court
Markham, Ontario L3R 8V2
Canada
Toll Free 800 227 7215
Tel 905 752 3300
Fax 905 752 3304

Asia/Pacific

Japan
Datex-Ohmeda K. K.
TRC Annex 9F
6-1-1 Heiwajima
Ohta-ku, Tokyo 143-0006
Japan
Tel 81 3 5763 6801
Fax 81 3 5763 6838

Datex-Ohmeda K. K.
Technical Center
TRC A Bldg. AE 4-8
6-1-1 Heiwajima
Ohta-ku, Tokyo 143-0006
Japan
Tel 81 3 5763 6850
Fax 81 3 5763 6852

Malaysia
Datex-Ohmeda
13 Jalan 223
Level 2 Bangunan O'connors
46100 Petaling Jaya
Selangor, Malaysia
Tel 60 3 754 7872
Fax 60 3 757 6948

Singapore
Datex-Ohmeda Pte. Ltd.
152 Beach Road
#12-05/07 Gateway East
Singapore 189721
Tel 65 391 8618
Fax 65 291 6618

Australia

Datex-Ohmeda Pty. Ltd.
Units 1 & 2
149 Arthur Street
P O Box 356
Homebush
NSW 2140
Australia
Tel 61 132 229
Fax 61 297 461796

Europe

France
Datex-Ohmeda S.A.S.
17 rue Jean-Elysée Dupuy
F-69410 Champagne Au Mont d'Or
France
Tel 33 01 30 68 60 00
Fax 33 04 78 43 26 58

Germany
Datex-Ohmeda GmbH
Dr-Alfred-Herrhausen-Allee 24
D-47228 Duisburg
Germany
Tel 49 2065 691 0
Fax 49 2065 691 236

Italy
Datex-Ohmeda S.p.A.
Via Cassanese, 100
20090 Segrate, Milan
Italy
Tel 39 2 21693431
Fax 39 2 26926226

Netherlands
Datex-Ohmeda B.V.
Kantemarsweg 18
Post Box 22
3870 CA Hoevelaken
Netherlands
Tel 31 33 253 5404
Fax 31 33 253 7223

Spain
Datex-Ohmeda S.L.
C/Manuel Tovar 26
28034 Madrid
Spain
Tel 34 1 334 26 00
Fax 34 1 358 12 84

United Kingdom
Datex-Ohmeda Ltd.
Ohmeda House
71 Great North Road
Hatfield Hertfordshire
AL9 5EN England
Tel 44 1707 263570
Fax 44 1707 260191

Latin America, Caribbean

Datex-Ohmeda Latin America
10685 North Kendall Drive
Miami, FL 33176
USA
Tel 1 305 273 9940
Fax 1 305 273 4382

For distributor locations worldwide
www.ohmedamedical.com

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Printed in USA
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